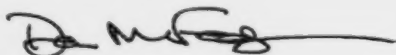# Directive 017

Revised edition October 22, 2009

Measurement Requirements for Upstream Oil and Gas Operations

The Energy Resources Conservation Board (ERCB/Board) has approved this directive on October 22, 2009.

Dan McFadyen
Chairman

Contents

Introduction

Purpose

Directive 017 consolidates, clarifies, and updates Energy Resources Conservation Board (ERCB) requirements related to the measurement of fluid (gas and liquid) production and disposition associated with upstream petroleum operations. The term "measurement" as used in this document means "measurement, accounting, and reporting." While measurement is the determination of a volume, accounting and reporting are integral components of measurement in that after a fluid volume is "measured," mathematical procedures (accounting) may have to be employed to arrive at the desired volume to be "reported."

What's New in This Edition

This revised edition of *Directive 017* includes two new sections:

- 12: Heavy Oil Measurement
- 13: Condensate and High Vapour Pressure Liquids Measurement

Highlights of other changes:

- Enforcement clause changed to refer reader to *Directive 019: ERCB Compliance Assurance—Enforcement.*
- Section 2.5.2.1: Item 9 has been added to provide another alternative to primary measurement element inspection.
- Section 4.2.4: Clarifies gas plant inlet measurement.
- Section 4.3.1: Absolute pressure transmitter is required for any vacuum measurement.
- Section 4.3.3.1: Reported total flare volume must exclude any fuel volumes.
- Section 4.3.3.2(a): All existing gas electronic flow measurement (EFM) requirements in *Directive 046: Production Audit Handbook* have been transferred to this section.
- Section 4.3.5.2: Clarification on GOR and GIS applications if criteria are met, no specific approval is required.
- Section 4.3.5.3: Produced gas measurement with heavy oil/crude bitumen has been changed to refer to Section 12.3.2
- Section 6.2.2.1: Clarifies Petroleum Registry of Alberta battery code requirements for oil well tests.
- Section 6.4.4.3: Field header well identification requirement has been transferred from *Directive 046.*
- Section 6.8: All existing oil EFM requirements in *Directive 046* have been transferred to this section.
- Section 7.2.4: Clarifies use of "ABMC" at SE Alberta shallow gas batteries.
- Sections 7.3.2 and 7.4.2: New calculation procedures have been added and existing calculation procedures have been modified.
- Section 8.4.1: Clarifies exceptions on sampling to match Section 8.4.2.

In the version available on the ERCB Web site www.ercb.ca, all changes made since the previous edition (February 2009) appear in blue type.

Intent

This directive specifies

- what and how volumes must be measured,
- what, where, and how volumes may be estimated,
- if accounting procedures must be performed on the determined volumes and what they are,
- what data must be kept for audit purposes, and
- what resultant volumes must be reported to the ERCB.

The licensee must comply with all requirements set out in this directive.

In this directive, the term "must" indicates a requirement that must be followed. In some situations, a requirement may be subject to exceptions if specific conditions are met.

The term "should" indicates a recommendation that will not be subject to enforcement. However, the ERCB may direct the licensee in writing to implement changes to improve measurement accuracy, and this direction will become a condition of operation for that facility or facilities.

The directive does not include instructions on how the volumes must be reported to the ERCB (which are included in other ERCB documents, such as *Directive 007: Production Accounting Handbook*), but it does include some information on requirements regarding subtype, status, and code in accordance with those documents.

Note that other sections relating to measurement listed in Appendix 1 are under development and will be added to the directive as they are completed.

If requirements in previously issued ERCB documents (interim directives, informational letters, guides, etc.) conflict with the requirements in this directive, the requirements in this directive replace the prior requirements. Over time, it is intended that all relevant superseded requirements will be rescinded.

ERCB documents fully or partially replaced by this directive are listed in the Appendix 2.

Definitions

Many terms used in this directive are defined in the Glossary (Appendix 3). However, many critically important definitions are also included within applicable sections.

Enforcement

The ERCB enforcement process is specified in *Directive 019: ERCB Compliance Assurance—Enforcement*. Noncompliance events are listed in the Risk Assessed Noncompliance section of *Directive 019* under the Operations Group, Production Operations Section, for *Directive 017*, *Directive 004: Determination of Water Production at Gas Wells*, and *Directive 007* requirements.

1 Standards of Accuracy

1.1 Introduction

With regard to accuracy, it is assumed an exact or true value exists for any variable that is valid for the conditions existing at the moment the result is determined. Determining the true value without doubt cannot be done, due to the limitations of measuring equipment and procedures and the possibility of human error. Typically, the closer one wants to approach the true value, the more expense and efforts have to be expended.

The ERCB has developed standards of accuracy for gas and liquid measurement that take into account such concerns as royalty, equity, reservoir engineering, declining production rates, and aging equipment. These standards have evolved, but originated from a 1972 Board hearing decision that determined a need for pool production accuracy standards of 2% for oil, 3% for gas, and 5% for water. The current standards are stated as "maximum uncertainty of monthly volume" and/or "single point measurement uncertainty." The uncertainties are to be applied as "plus/minus" (e.g., ± 5%). Measurement at delivery/sales points must meet the highest accuracy standards because volumes determined at these points have a direct impact on royalty determination. Other measurement points that play a role in the overall accounting process are subject to less stringent accuracy standards to accommodate physical limitations and/or economics.

The specific standards of accuracy are listed in Section 1.8.

1.2 Applicability and Use of Uncertainties

The ERCB used the uncertainty levels contained in this section to develop many of the requirements for equipment and/or procedures relating to measurement, accounting, and reporting for various aspects of oil and gas production and processing operations, which are explained in detail in other sections. **If those requirements are being met, the ERCB considers a licensee to be in compliance without the need to demonstrate compliance with the applicable uncertainty requirements contained in this section. In other words, licensees need not have any concern about the stated uncertainty limits for operations that meet minimum requirements, exception requirements, or specific ERCB approval requirements as set out in this directive. ERCB inspectors and auditors do not look for compliance with the stated uncertainty limits in conducting inspections and audits.**

In some cases a licensee may wish to deviate from the minimum requirements for equipment and/or procedures that are stated in this guide. That may be acceptable if it is in accordance with the following:

- A licensee may deviate from the minimum requirements without specific ERCB approval if no royalty, equity, or reservoir engineering concerns are associated with the volumes being measured and the licensee is able to demonstrate that the alternative measurement equipment and/or procedures will provide measurement accuracy within the applicable uncertainties. In some cases, as described in Section 5: Site-specific Deviation from Base Requirements, the licensee does not need to demonstrate compliance with the applicable uncertainties, but may instead demonstrate compliance with other specific criteria. In such cases, ERCB inspectors and auditors will review the licensees' records for demonstrated compliance with the uncertainty limits or with the other specified criteria.

- If royalty, equity, or engineering concerns are associated with the volumes being measured, a licensee may be allowed, upon application to the Operations Group of the

ERCB's Compliance and Operations Branch, to deviate from the minimum requirements. The application must demonstrate that the proposed alternative measurement equipment and/or procedures will either provide measurement accuracy within the applicable uncertainties or meet specific criteria described in Section 5: Site-specific Deviation from Base Requirements. Applications will also be considered if measurement accuracy will be marginally outside the uncertainty limits or specified criteria will be marginally exceeded. In such cases, ERCB inspectors and auditors will review the licensees' records for documentation to confirm that approval has been obtained to deviate from the minimum requirements and for compliance with the approval conditions.

Note that crude bitumen wells and heavy oil wells (density of 920 kilograms per cubic metre [kg/m^3] or greater at 15°C) have specific measurement requirements that in some cases differ from those applicable to conventional oil wells. In those cases, the crude bitumen and heavy oil measurement requirements are based on operational characteristics and limitations, rather than uncertainty limits. The uncertainties described in this section do not apply to measurement of heavy oil production associated with the following oil systems:

- total battery gas
- well oil (proration battery)
- well gas (proration battery)
- well water (proration battery)

1.3 Maximum Uncertainty of Monthly Volume

The ERCB requires production data to be reported on a calendar month basis. "Maximum uncertainty of monthly volume" relates to the limits applicable to equipment and/or procedures used to determine the total monthly volume. Total monthly volumes may result from a single month-long measurement, but more often result from a combination of individual measurements and/or estimations. For example, consider a well in an oil proration battery to which a maximum uncertainty of monthly volume would apply:

- First, the well is tested, and the oil test rate is used to estimate the well's production for the period until the next test is conducted.

- The well's total estimated oil production for the month is combined with the month's estimated oil production for the other wells in the battery to arrive at the total estimated monthly oil production for the battery.

- The total actual monthly oil production for the battery is determined based on measured deliveries out of the battery and inventory change.

- A proration factor is determined by dividing the actual battery production by the estimated battery production.

- The proration factor is multiplied by the well's estimated production to determine the well's actual monthly production.

1.4 Single Point Measurement Uncertainty

"Single point measurement uncertainty" relates to the limits applicable to equipment and/or procedures used to determine a single-phase specific volume at a single measurement point. The oil volume determined during a 24-hour well test conducted on a well in a proration

battery is an example of a specific volume determination to which a single point measurement uncertainty limit would apply.

1.5 Confidence Level

The stated uncertainties are not absolute limits. The confidence level, which indicates the probability that true values will be within the stated range, is 95%. This implies that there is a 95% probability (or 19 chances in 20) that the true value will be within the stated range.

1.6 Determination of Uncertainties

The uncertainties referred to relate to the accuracies associated with measurement devices, device calibration, sample gathering and analysis, variable operating conditions, etc. These uncertainties are for single-phase specific volume determination points of specific fluids (oil, gas, or water) or for combinations of two or more such points. These uncertainties do not relate to comparisons of two or more measurement points, such as comparison of inlet volumes to outlet volumes. Such comparisons are typically expressed as proration factors, allocation factors, or metering differences.

The uncertainties are relevant to equipment at the time of installation. No uncertainty adjustment is required to account for the effects of multiphase fluids, wear, sludge or scale buildup, etc., as it is accepted that such conditions would constitute a bias error to be monitored and accounted for through the use of proration factors, allocation factors, or metering differences.

The methods to be used for determining and combining uncertainties are found in the latest edition of the American Petroleum Institute (API) *Manual of Petroleum Measurement Standards* (MPMS), Chapter 13: Statistical Aspects of Measuring and Sampling or the latest edition of the International Standard Organization (ISO) Standard 5168: Measurement of Fluid Flow—Estimation of Uncertainty of a Flow-rate Measurement.

1.6.1 Example Calculation

Determination of single point measurement uncertainty for well oil (proration battery) using "root sum square" methodology:

Individual uncertainties from historical ERCB research:

For oil/emulsion measurement,
Oil meter uncertainty = 0.5% (typical manufacturer's specification)
Meter proving uncertainty = 1.5%
Sediments and water (S&W) determination uncertainty = 0.5%
Combined uncertainty $= \sqrt{[(0.5)^2 + (1.5)^2 + (0.5)^2]}$
$= 1.66\%$ (rounded to 2.0%)
For delivery point gas measurement,
Primary measurement device – gas meter uncertainty = 1.0%
Secondary device – (pulse counter or transducer, etc.) uncertainty = 0.5%
Secondary device calibration uncertainty = 0.5%
Tertiary device – (flow calculation, EFM, etc.) uncertainty = 0.2%
Gas sampling and analysis uncertainty = 1.5%
Combined uncertainty $= \sqrt{[(1.0)^2 + (0.5)^2 + (0.5)^2 + (0.2)^2 + (1.5)^2]}$
$= 1.95\%$ (rounded to 2.0%)

1.7 Explanation of Standards of Accuracy

1.7.1 Oil Systems

(i) Total battery/facility oil (delivery point measurement), including single-well batteries

For Figure 1.1,

> **m** = single point measurement uncertainty
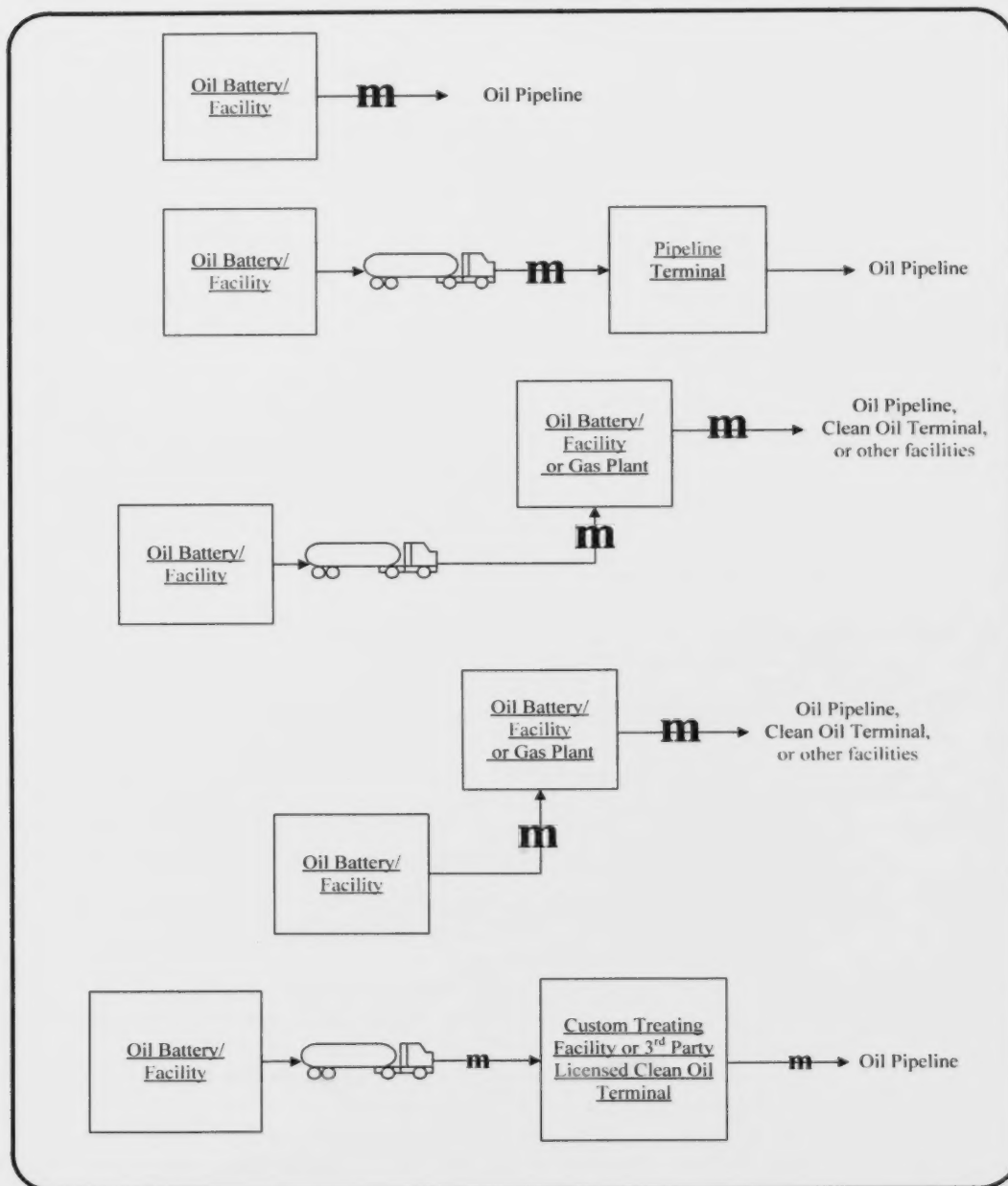
Figure 1.1. Total battery/facility oil (delivery point measurement)

Maximum uncertainty of monthly volume = N/A
The uncertainty of the monthly volume will vary, depending upon the number of individual measurements that are combined to yield the total monthly volume.

Single point measurement uncertainty:
Delivery point measures > 100 m^3/d = 0.5%
Delivery point measures ≤ 100 m^3/d = 1%

(ii) **Total battery gas** (includes produced gas that is vented, flared, or used as fuel), including single-well batteries—also referred to as "associated gas," as it is the gas produced in association with oil production at oil wells

For Figure 1.2,
(**m**) = single point measurement uncertainty
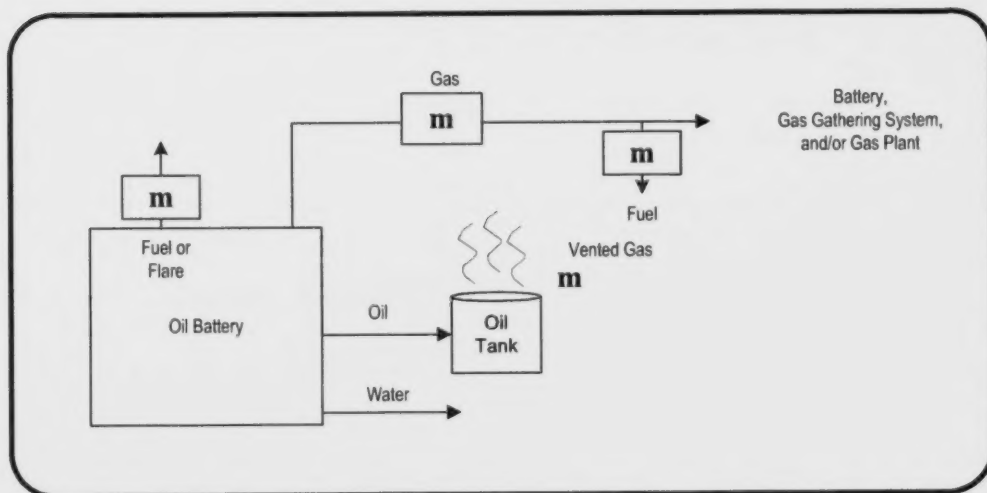
Figure 1.2

Single point measurement uncertainty:
> 16.9 10^3 m^3/d = 3%
> 0.50 10^3 m^3/d but ≤ 16.9 10^3 m^3/d = 3%
≤ 0.50 10^3 m^3/d = 10%

Maximum uncertainty of monthly volume (**M**)
> 16.9 10^3 m^3/d = 5%
> 0.50 10^3 m^3/d but ≤ 16.9 10^3 m^3/d = 10%
≤ 0.50 10^3 m^3/d = 20%
Note that **M** is dependent upon combined deliveries, fuel, and vented gas measurement.

The maximum uncertainty of total monthly battery gas volumes allows for reduced emphasis on accuracy as gas production rate declines. For gas rates up to 0.50 10^3 m^3/d, the gas volumes may be determined by using estimates; therefore, the maximum uncertainty of monthly volume is set at 20%. If gas rates exceed 0.50 10^3 m^3/d, the gas must be measured; however, a component of the total monthly gas volume may include estimates for low volumes of fuel, vented, or flare gas that may add to the monthly uncertainty. At the highest gas production rates, it is expected the use of estimates will be minimal or at least have a

minor impact on the accuracy of the total monthly gas volume, thereby resulting in the 5% maximum uncertainty of monthly volume.

The equipment and/or procedures used to determine the measured gas volumes (when measurement is required) must be capable of meeting a 3% single point measurement uncertainty. Due to the difficulty associated with measuring very low gas rates, the equipment and/or procedures used in determining gas-oil ratios or other factors to be used in estimating gas volumes where rates do not exceed 0.50×10^3 m³/d are expected to be capable of meeting a 10% single point measurement uncertainty.

These uncertainties do not apply to gas produced in association with heavy oil (density of 920 kg/m³ or greater at 15°C).

(iii)   **Total battery water,** including single-well batteries

For Figure 1.3,
M = maximum uncertainty of monthly volume
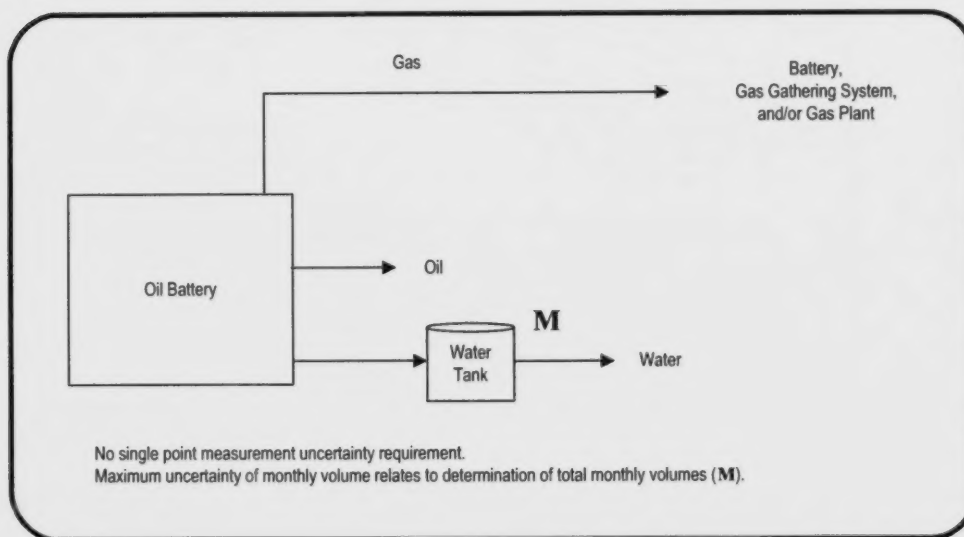
Figure 1.3

Maximum uncertainty of monthly volume:
> 50 m³/month = 5%
≤ 50 m³/month = 20%

Single point measurement uncertainty = N/A
Total battery water may be determined by measurement or estimation, depending on production rates, so no basic requirement has been set for single point measurement uncertainty.

Total battery water production volumes not exceeding 50 m³/month may be determined by estimation; therefore, the maximum uncertainty of monthly volume is set at 20%.

If the total battery water production volumes exceed 50 m^3/month, the water must be separated from the oil and measured; therefore, the maximum uncertainty of monthly volume is set at 5%.

(iv) Well oil (proration battery)

For Figure 1.4,
(m) = single point measurement uncertainty
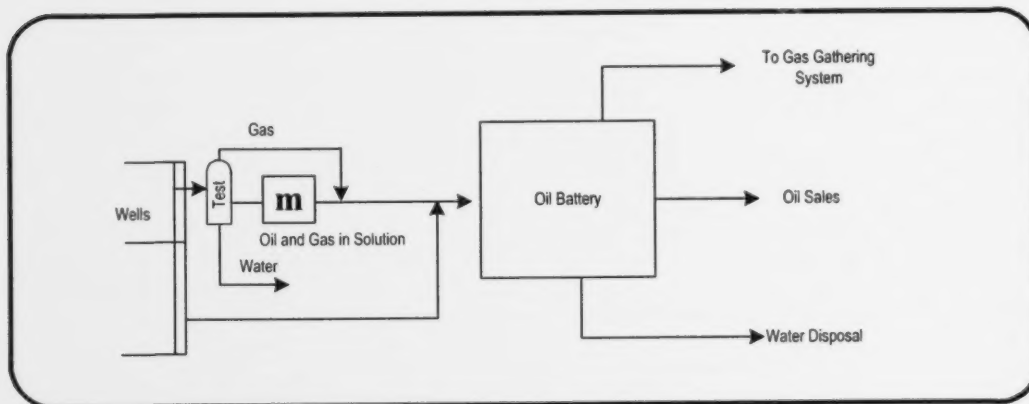
Figure 1.4

Single point measurement uncertainty:
All classes = 2%

Maximum uncertainty of monthly volume (**M**):
Class 1 (high) > 30 m^3/d = 5%
Class 2 (medium) > 6 m^3/d but ≤ 30 m^3/d = 10%
Class 3 (low) > 2 m^3/d but ≤ 6 m^3/d = 20%
Class 4 (stripper) ≤ 2 m^3/d = 40%
M is dependent upon oil and gas test volumes and the number of days the test is used for estimating production, plus correction by a proration factor.

The maximum uncertainty of monthly well oil production volumes for light and medium density oil wells in proration batteries has been developed to allow for reduced emphasis on accuracy as oil production rates decline. Rather than being determined by continuous measurement, monthly well oil production volumes are estimated from well tests and corrected by the use of proration factors to result in "actual" volumes. Lower rate wells are allowed reduced testing frequencies, which, coupled with the fact that wells may exhibit erratic production rates between tests, results in less certainty that the reported monthly oil production volume will be accurate.

The equipment and/or procedures used to determine oil volumes during the well tests must be capable of meeting a 2% single point measurement uncertainty for all classes of wells.

These uncertainties do not apply to heavy oil wells (density of 920 kg/m^3 or greater at 15°C) in proration batteries.

(v) **Well gas (proration battery)**—also referred to as "associated gas," as it is the gas produced in association with oil production at oil wells

For Figure 1.5,
(**m**) = single point measurement uncertainty
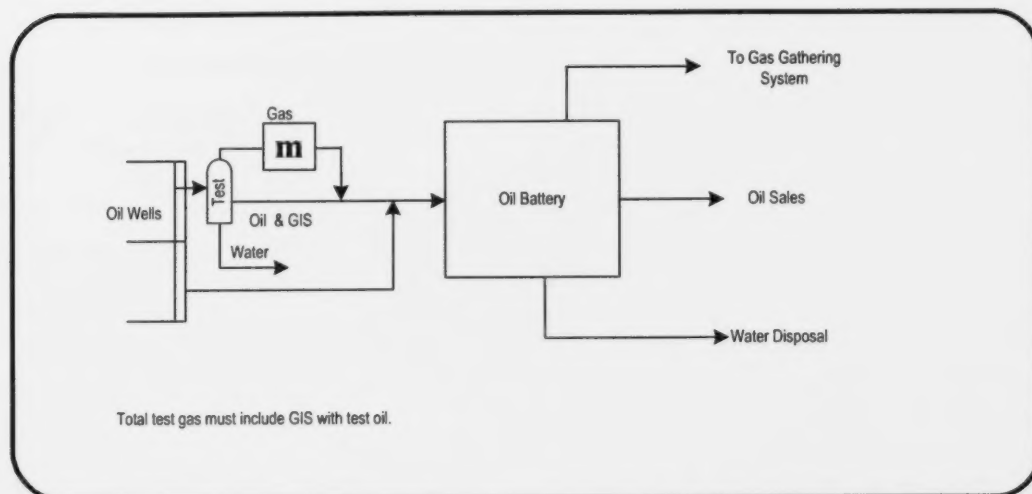
Figure 1.5

Single point measurement uncertainty:
> $16.9 \ 10^3 \ m^3/d = 3\%$
> $0.50 \ 10^3 \ m^3/d$ but $\leq 16.9 \ 10^3 \ m^3/d = 3\%$
$\leq 0.50 \ 10^3 \ m^3/d = 10\%$

Maximum uncertainty of monthly volume (**M**):
> $16.9 \ 10^3 \ m^3/d = 5\%$
> $0.50 \ 10^3 \ m^3/d$ but $\leq 16.9 \ 10^3 \ m^3/d = 10\%$
$\leq 0.50 \ 10^3 \ m^3/d = 20\%$

M is dependent upon oil and gas test volumes and the number of days the test is used for estimating production, plus correction by a proration factor.

The maximum uncertainty of monthly oil well gas volumes has been developed to allow for reduced emphasis on accuracy as gas production rates decline. Rather than being determined by continuous measurement, monthly oil well gas production volumes are estimated from well tests and corrected by the use of proration factors to result in "actual" volumes. Low gas production rates are typically associated with wells that are allowed reduced testing frequencies, which, coupled with the fact that wells may exhibit erratic production rates between tests, results in less certainty that the reported monthly gas production volume will be accurate.

For gas rates up to $0.50 \ 10^3 \ m^3/d$, the well test gas volume may be determined by using estimates; therefore, the maximum uncertainty of monthly volume is set at 20%. If gas rates exceed $0.50 \ 10^3 \ m^3/d$, the test gas must be measured; however, a component of a well's total test gas volume may include estimates for solution gas dissolved in the test oil volume (gas-in-solution), which may add to the monthly uncertainty. At the highest gas production rates, it

is expected that the use of estimates will be minimal or at least have a minor impact on the accuracy of the total monthly gas volume, thereby resulting in the 5% maximum uncertainty of monthly volume.

The equipment and/or procedures used to determine the measured test gas volumes (if measurement is required) must be capable of meeting a 3% single point measurement uncertainty. Due to the difficulty associated with measuring very low gas rates, the equipment and/or procedures used in determining gas-oil ratios or other factors to be used in estimating gas volumes if rates do not exceed $0.50 \ 10^3 \ m^3/d$ are expected to be capable of meeting a 10% single point measurement uncertainty.

These uncertainties do not apply to gas produced by heavy oil wells (density of 920 kg/m^3 or greater at 15°C) in proration batteries.

(vi) Well water (proration battery)

For Figure 1.6,
(**m**) = single point measurement uncertainty
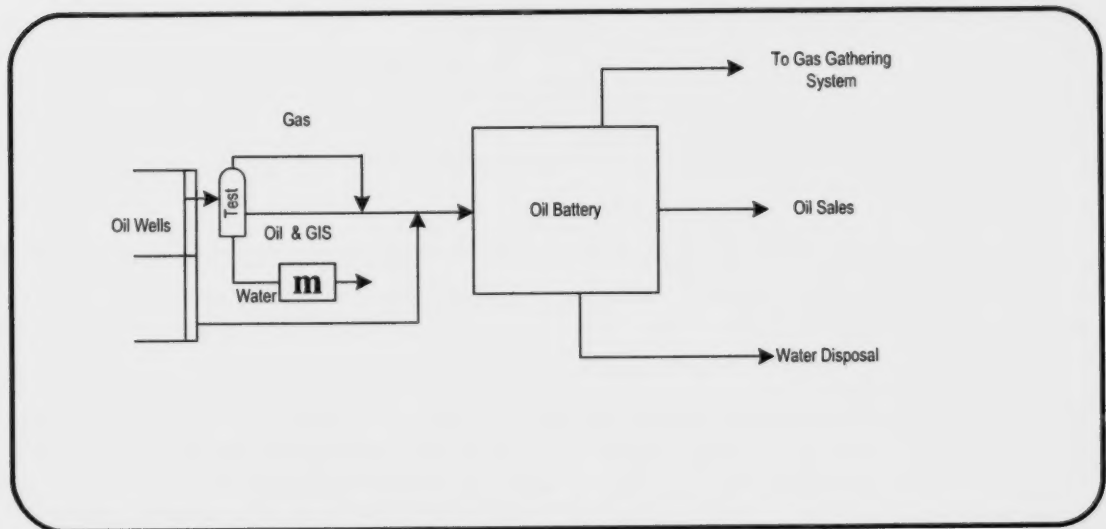
Figure 1.6

Single point measurement uncertainty = 10%

Maximum uncertainty of monthly volume = N/A
The uncertainty of the monthly volume will vary, depending upon the method used to determine test water rates and the frequency of well tests.

Rather than being determined by continuous measurement, monthly oil well water production volumes are estimated from well tests and corrected by the use of proration factors to result in "actual" volumes. The water rates determined during the well tests may be inferred from determining the water content of emulsion samples, and in some cases estimates may be used to determine water rates. Therefore, the single point measurement uncertainty is set at 10%.

These uncertainties do not apply to heavy oil wells (density of 920 kg/m^3 or greater at 15°C) in proration batteries.

1.7.2 Gas Systems

(i) Gas deliveries (sales gas)

For Figure 1.7,
(**m**) = single point measurement uncertainty
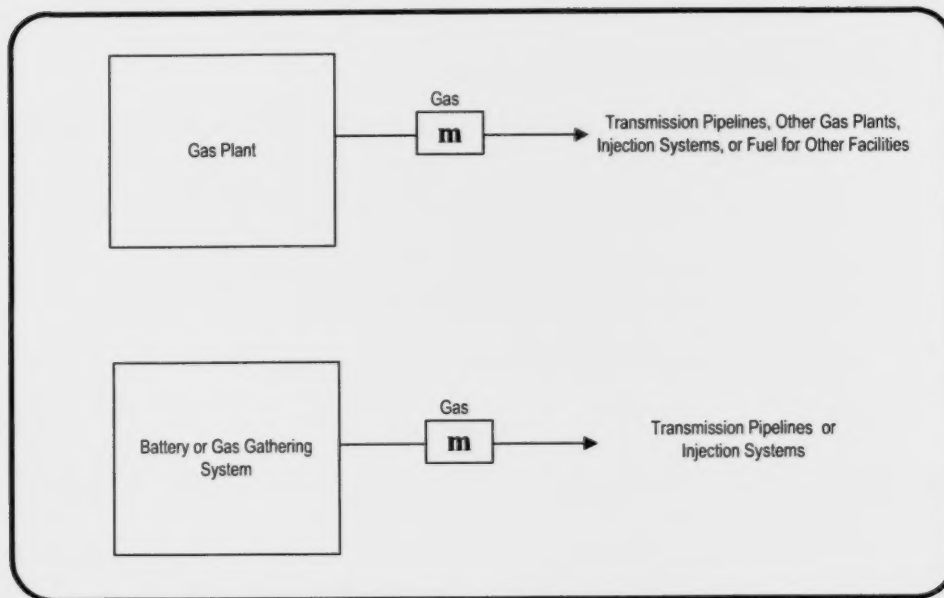
Figure 1.7

Single point measurement uncertainty = 2%

Maximum uncertainty of monthly volume = N/A
The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

Gas deliveries in this context will typically be clean, processed sales gas that is delivered out of a gas plant into a transmission pipeline. The measurement at this point determines the gas volumes upon which royalties will be based. Therefore, a stringent expectation is set for the single point measurement uncertainty. In some cases, this type of gas may be delivered to other plants for further processing or to injection facilities.

(ii) Hydrocarbon liquid deliveries

For Figure 1.8,
(**m**) = single point measurement uncertainty
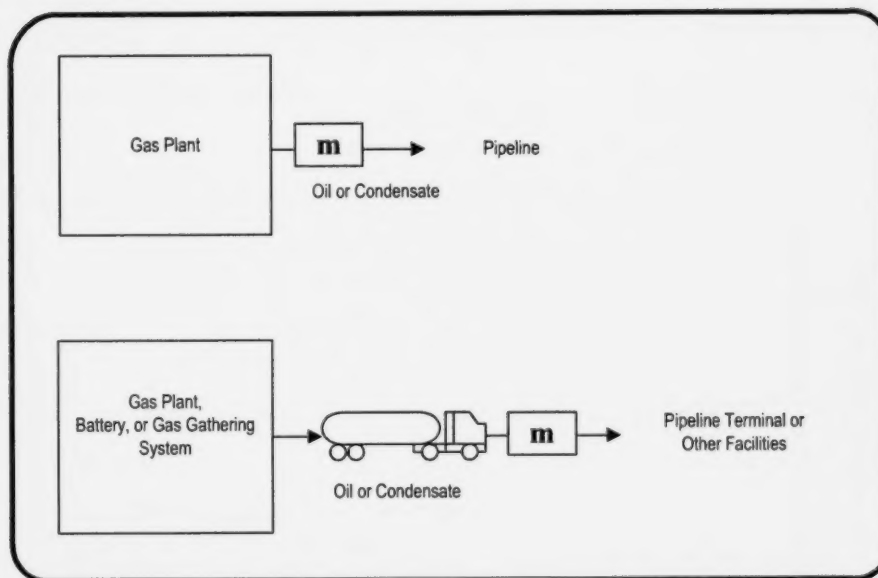
Figure 1.8

Single point measurement uncertainty:
Delivery point measures > 100 m^3/d = 0.5%
Delivery point measures ≤ 100 m^3/d = 1%

Maximum uncertainty of monthly volume = N/A
The uncertainty of the monthly volume will vary, depending upon the number of individual measurements that are combined to yield the total monthly volume.

The term "delivery point measurement" for hydrocarbon liquids refers to the point at which the hydrocarbon liquid production from a battery or facility is measured. Where clean hydrocarbon liquids are delivered directly into a pipeline system (Lease Automatic Custody Transfer [LACT] measurement) or trucked to a pipeline terminal, it can also be referred to as the "custody transfer point." The "delivery point" terminology is from the perspective of the producing battery or facility, but the receiving facility (pipeline, terminal, custom treating facility, other battery, etc.) may refer to this point as its "receipt point." The hydrocarbon liquid volume determined at the delivery point is used in all subsequent transactions involving that liquid.

Hydrocarbon liquids delivered out of a gas system at the well, battery, or plant inlet level are typically condensate, and in some cases they may be considered to be oil. The hydrocarbon liquids delivered out of a gas plant may be pentanes plus, butane, propane, ethane, or a mixture of various components. The volumes determined at this point are the volumes upon which royalties are based.

The measurement equipment and/or procedures must be capable of determining the hydrocarbon liquid volume within the stated limits.

For facilities where the hydrocarbon liquid delivery volumes total ≤ 100 m³/d, the single point measurement uncertainty has been increased to allow for the economical handling of hydrocarbon liquids when minimal volumes would not justify the added expense for improved measurement equipment and/or procedures.

(iii) Plant inlet or total battery/group gas

For Figure 1.9,
M = maximum uncertainty of monthly volume
(m) = single point measurement uncertainty

M is dependent upon combined uncertainties of measured gas and gas equivalent of condensate.

M is dependent upon combined uncertainties of measured gas and gas equivalent of recombined condensate.

M is dependent upon uncertainties of measured gas only.
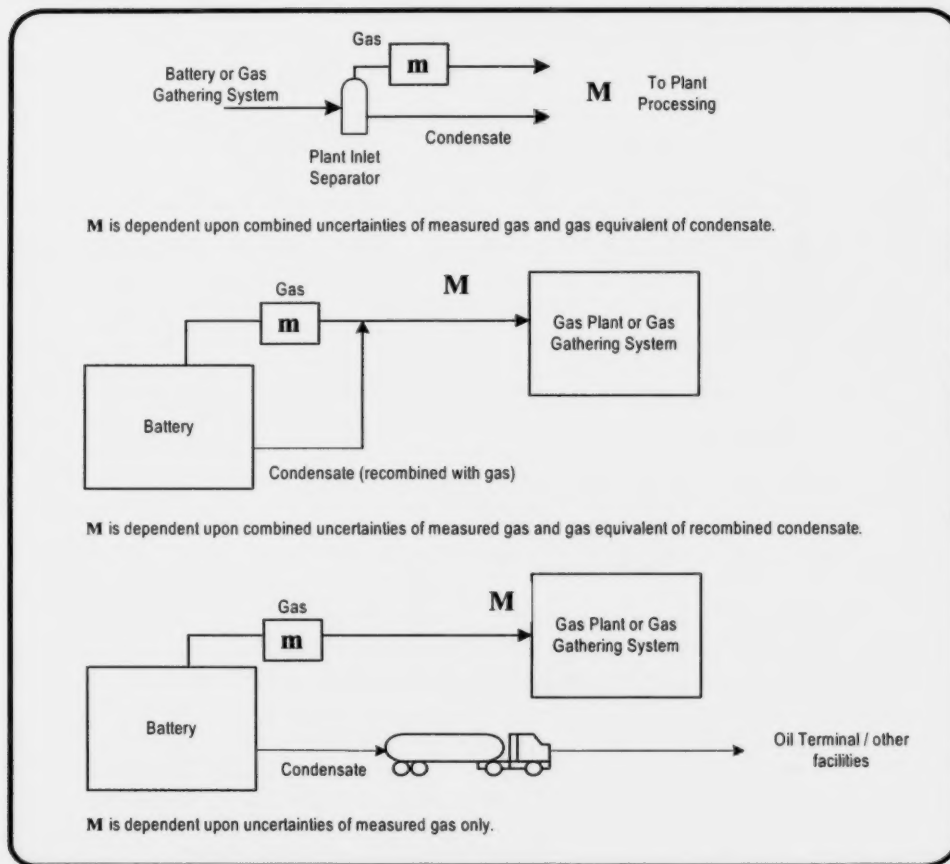
Figure 1.9

Maximum uncertainty of monthly volume = 5%

Single point measurement uncertainty = 3%

Plant inlet gas or total battery/group gas is typically unprocessed gas that may vary in composition and may contain entrained liquids. The total reported gas volume could result from combining several measured volumes from various points and may also include the calculated gas equivalent volume of entrained hydrocarbon liquids (typically condensate). The expectation for the maximum uncertainty of monthly volume is set at 5% to allow for the uncertainties associated with measuring gas under those conditions.

The equipment and/or procedures used to determine the measured gas volumes must be capable of meeting a 3% single point measurement uncertainty.

(iv) Plant inlet or total battery/group condensate (recombined)

For Figure 1.10,
(**m**) = single point measurement uncertainty
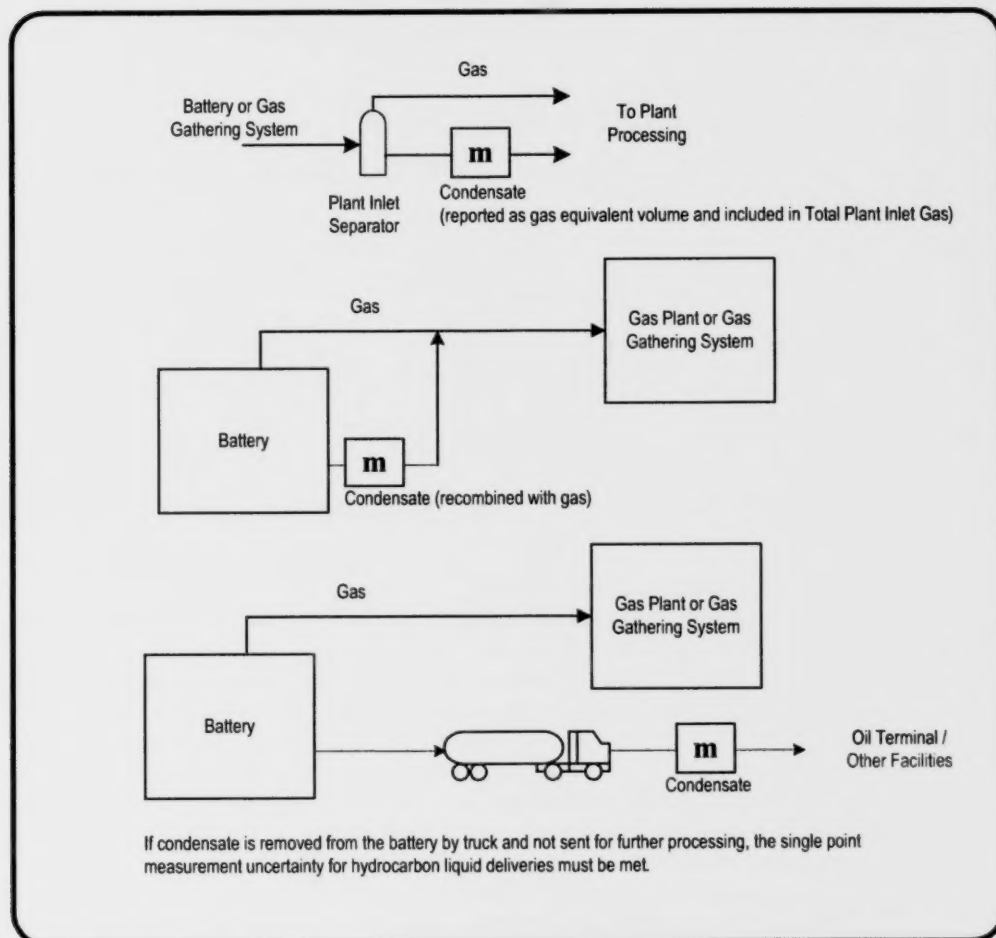
Figure 1.10

Single point measurement uncertainty = 2%

Maximum uncertainty of monthly volume = N/A
The condensate volume is included in the total gas volume for reporting purposes and is therefore covered by the maximum uncertainty of monthly volume for the plant inlet or total battery/group gas.

Plant inlet condensate is typically separated from the inlet stream and sent through the plant for further processing. For reporting purposes, the gas equivalent of the plant inlet condensate is included in the total plant inlet gas volume. If total battery/group condensate upstream of the plant inlet is separated and measured prior to being recombined with the gas production,

the condensate is converted to a gas equivalent volume and included in the gas production volume. In either case, the condensate single point measurement uncertainty is set at 2% for the liquid volume determination.

Note that if plant inlet or total battery/group condensate is separated and delivered out of the system at that point, the condensate measurement is subject to the single point measurement uncertainties stipulated for hydrocarbon liquid deliveries (above).

(v) Fuel gas

For Figure 1.11,
(**m**) = single point measurement uncertainty
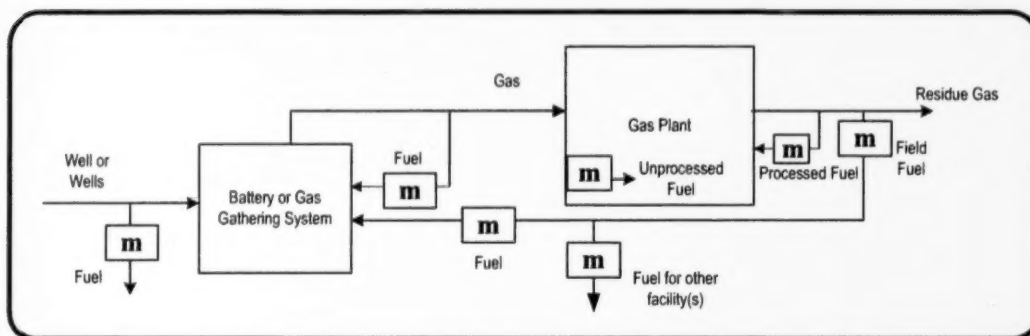
Figure 1.11

Single point measurement uncertainty:
$> 0.50 \ 10^3 \ m^3/d = 3\%$
$\leq 0.50 \ 10^3 \ m^3/d = 10\%$

Maximum uncertainty of monthly volume (**M**):
$> 0.50 \ 10^3 \ m^3/d = 5\%$
$\leq 0.50 \ 10^3 \ m^3/d = 20\%$

Note that **M** is dependent upon combined uncertainties of various fuel sources at each reporting facility.

The maximum uncertainty of monthly fuel gas volumes allow for reduced emphasis on accuracy as gas flow rates decline.

For all upstream oil and gas facilities, if the annual average fuel gas rate is $0.50 \ 10^3 \ m^3/d$ or less on a per-site basis, the gas volume may be determined by using estimates. Therefore, the maximum uncertainty of the monthly volume is set at 20%. If the annual average fuel gas rates exceed $0.50 \ 10^3 \ m^3/d$ on any site, the gas must be measured, but since the gas being used as fuel may be unprocessed gas and part of the total fuel gas volume may include some estimated volumes (up to $0.50 \ 10^3 \ m^3/d$), the maximum uncertainty of the monthly volume is set at 5% to allow for the uncertainties associated with measuring gas under those conditions.

The equipment and/or procedures used to determine the measured gas volumes (if measurement is required) must be capable of meeting a 3% single point measurement uncertainty. Due to the difficulty associated with measuring very low gas rates, the equipment and/or procedures used in determining gas-oil ratios or other factors to be used in estimating

gas volumes if rates do not exceed 0.50 10^3 m^3/d are expected to be capable of meeting a 10% single point measurement uncertainty.

(vi) Flare gas

For Figure 1.12,
M = maximum uncertainty of monthly volume
(m) = single point measurement uncertainty
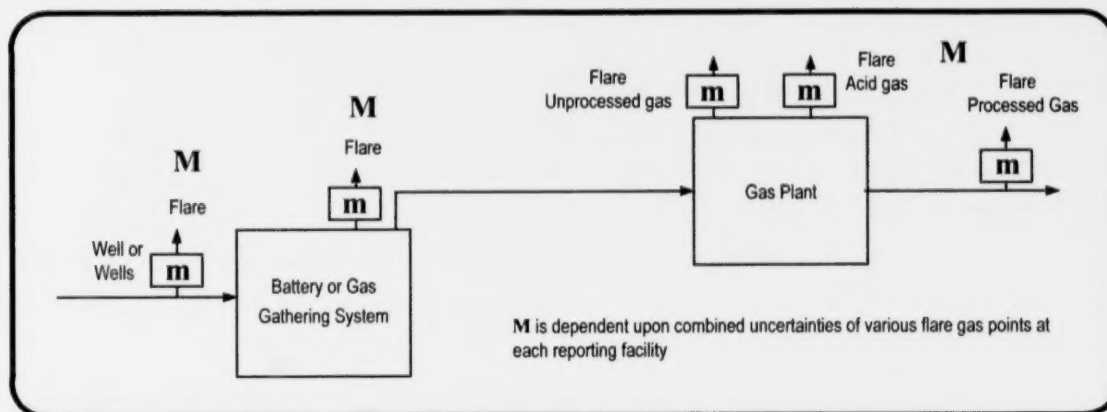
Figure 1.12

Maximum uncertainty of monthly volume = 20%

Single point measurement uncertainty = 5%

Flare gas may be clean processed gas or it may be unprocessed gas, depending on the point in the system from which gas is being flared. Continuous or intermittent flare and vent sources at all oil and gas production and processing facilities (excluding heavy oil and crude bitumen) where annual average total flared and vented volumes per facility exceed 0.5 10^3 m^3/d (excluding pilot, purge, or dilution gas) must be measured. Flare lines usually operate in a shut-in condition and may be required to accommodate partial or full volumes of gas production during flaring conditions. In some cases if flaring is infrequent and no measurement equipment is in place, flare volumes must be estimated (such as flaring at southeastern Alberta gas wells in a proration battery where there is no on-site measurement equipment). Therefore, the maximum uncertainty of the monthly volume is set at 20%, to allow for the erratic conditions associated with flare measurement.

The equipment and/or procedures used to determine the measured gas volumes (if measurement, not an estimate, is required) must be capable of meeting a 5% single point measurement uncertainty.

(vii) Acid gas

For Figure 1.13,
(m) = single point measurement uncertainty
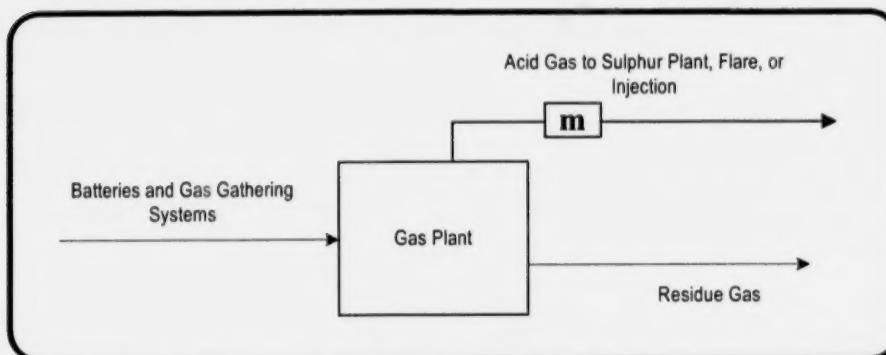
Figure 1.13

Single point measurement uncertainty = 10%

Maximum uncertainty of monthly volume = N/A
The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

Acid gas usually contains a great deal of water vapour and has other conditions associated with it, such as very low pressure that affects measurement accuracy. Therefore, the single point measurement uncertainty is set at 10%.

(viii) Dilution gas

For Figure 1.14,
M = maximum uncertainty of monthly volume
(m) = single point measurement uncertainty

M is dependent upon combined uncertainties of various dilution gas sources.
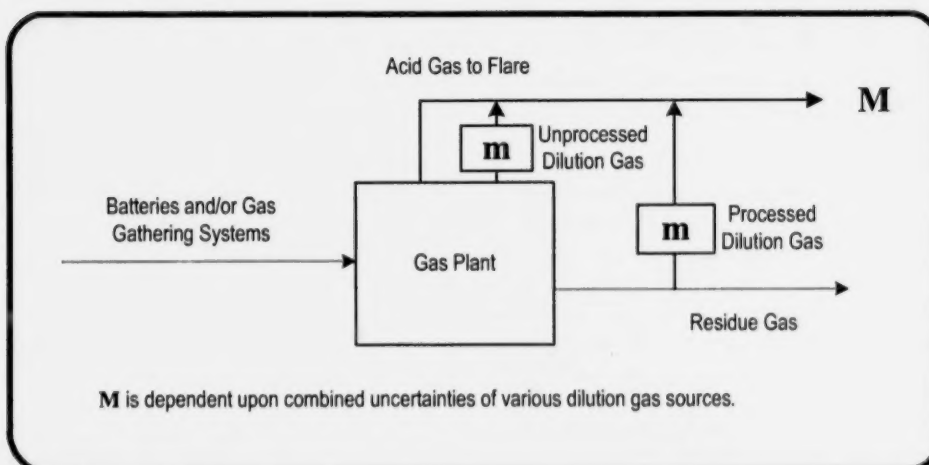
Figure 1.14

Maximum uncertainty of monthly volume = 5%

Single point measurement uncertainty = 3%

Dilution gas is typically "fuel" gas used to provide adequate fuel for incineration or flaring of acid gas. Since it must be measured, it is subject to the same uncertainties as stated above for fuel gas that must be determined by measurement.

(ix) Well gas (well site separation)

For Figure 1.15,
M = maximum uncertainty of monthly volume
(m) = single point measurement uncertainty
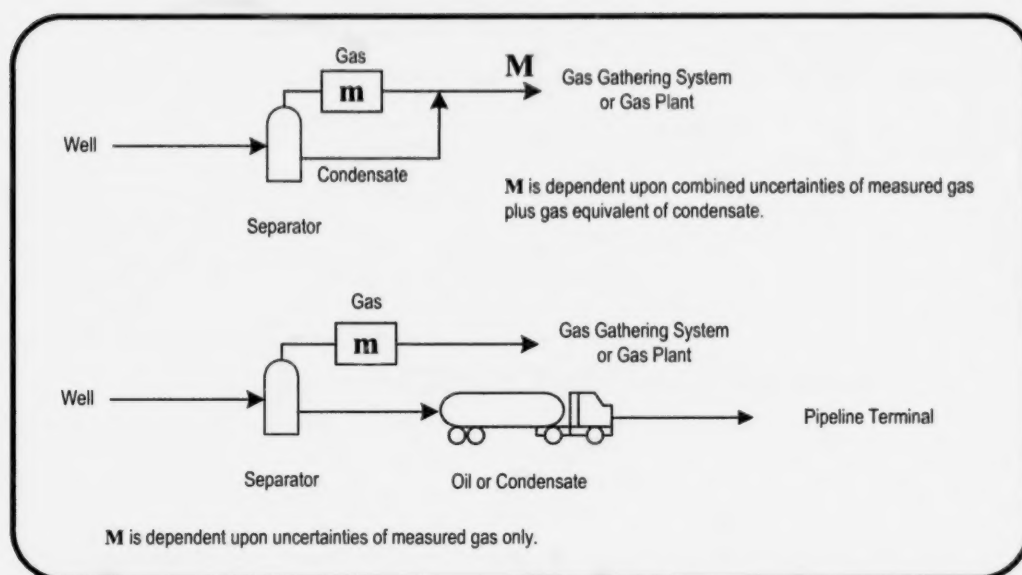
Figure 1.15

Maximum uncertainty of monthly volume:
$> 16.9 \times 10^3 \text{ m}^3/\text{d} = 5\%$
$\leq 16.9 \times 10^3 \text{ m}^3/\text{d} = 10\%$

Single point measurement uncertainty = 3%

If production components from gas wells are separated and continuously measured, the maximum uncertainty of monthly well gas volumes allows for reduced emphasis on accuracy as gas production rates decline. Since the separated gas is unprocessed and may still contain entrained liquids at the measurement point and a component of the total reported well gas production may include the calculated gas equivalent volume of the well's condensate production, the maximum uncertainty of monthly volumes also allows for the uncertainties associated with measuring gas under those conditions.

The equipment and/or procedures used to determine the separated measured well gas volumes must be capable of meeting a 3% single point measurement uncertainty.

(x) Well gas (proration battery)

For Figures 1.16 and 1.17,
M = maximum uncertainty of monthly volume
(m) = single point measurement uncertainty
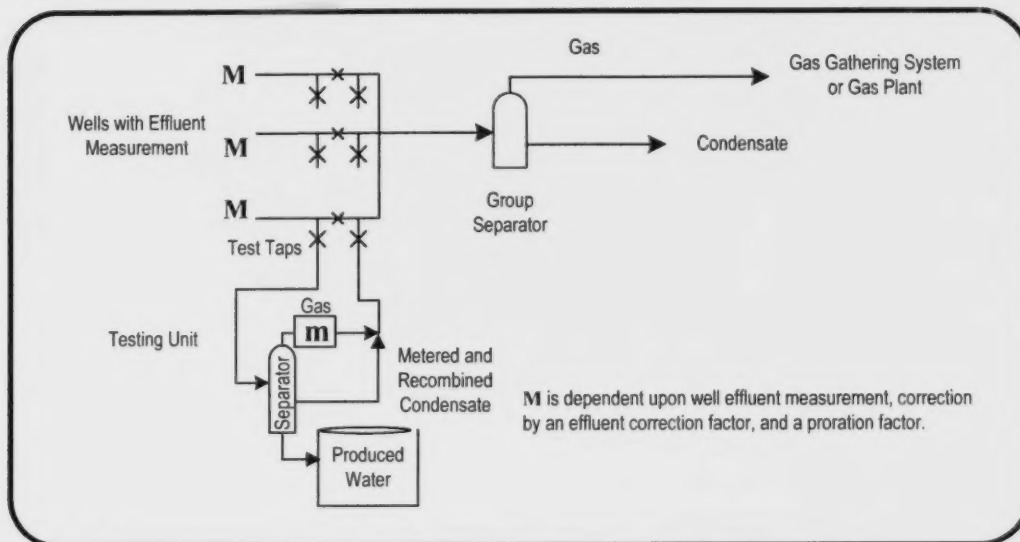
Figure 1.16. Well gas (effluent proration battery)
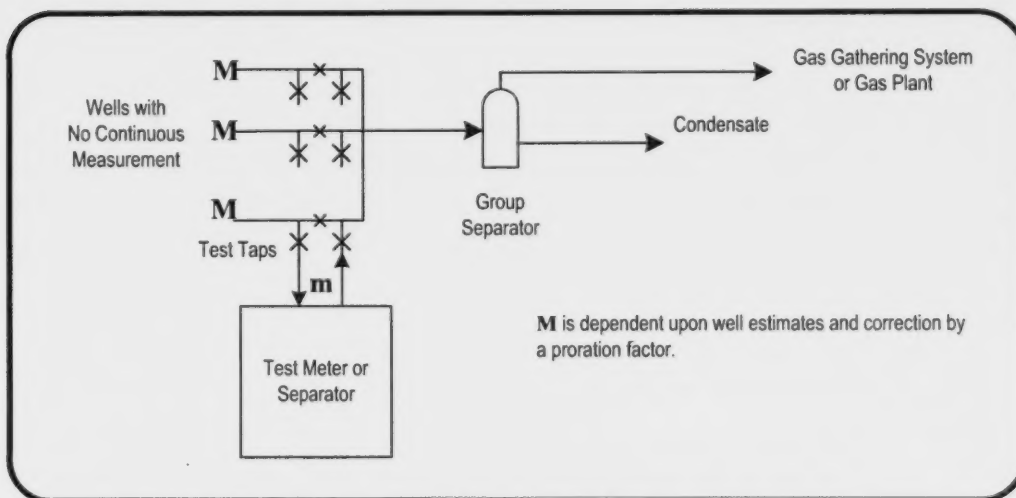
Figure 1.17. Well gas (Southeastern Alberta or other approved proration battery)

Maximum uncertainty of monthly volume = 15%

Single point measurement uncertainty = 3%

If production components from gas wells are not separated and continuously measured, the gas wells are subject to a proration accounting system. There are two types of gas proration batteries. "Wet" gas wells have continuous effluent measurement, and the "actual" production

is prorated based on the measurement of group gas and liquid components following separation at a central location. "Dry" gas wells approved to operate without continuous measurement have the production estimated based on periodic tests, and the actual production is prorated based on the measurement of group volumes at a central location. For both types of proration batteries, the maximum uncertainty of the monthly well gas volume is set at 15% to allow for the inaccuracies associated with these types of measurement systems.

The equipment and/or procedures used to determine the measured well test gas volumes downstream of separation during effluent meter correction factor tests or during the periodic dry gas well tests must be capable of meeting a 3% single point measurement uncertainty.

(xi) Well condensate (recombined)

For Figure 1.18,
(**m**) = single point measurement uncertainty
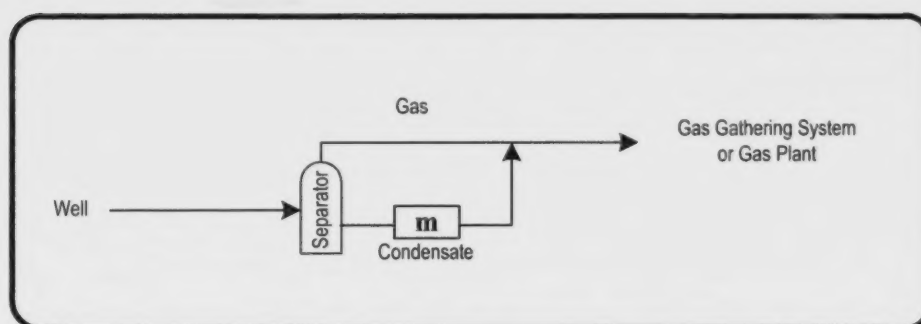
Figure 1.18

Single point measurement uncertainty = 2%

Maximum uncertainty of monthly volume = N/A
The gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes and is therefore covered by the monthly uncertainty for the well gas.

If condensate produced by a gas well is separated and measured at the wellhead prior to being recombined with the gas production, the condensate is mathematically converted to a gas equivalent volume and added to the well gas production volume. In this case, the condensate single point measurement uncertainty is set at 2% for the liquid volume determination. No requirement has been set for the maximum uncertainty of monthly volume because the gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes.

In the case of a gas well subject to effluent measurement, the gas equivalent of the condensate volume is included in the well's total gas production volume. The liquid volume determination, which is done during the effluent meter correction factor test, is subject to a single point measurement uncertainty of 2%. No requirement has been set for the maximum uncertainty of monthly volume because the gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes.

Note that if condensate produced by a gas well is separated at the wellhead and delivered out of the system at that point, the condensate is reported as a liquid volume. In this case, the

condensate measurement is subject to the single point measurement uncertainties stipulated for hydrocarbon liquid deliveries (above).

(xii) Total battery water

For Figure 1.19,
M = maximum uncertainty of monthly volume

Total battery water may be the water separated at the battery, the sum of the water separated at the wells, or a combination. M is dependent upon the method used to determine the total water volume.
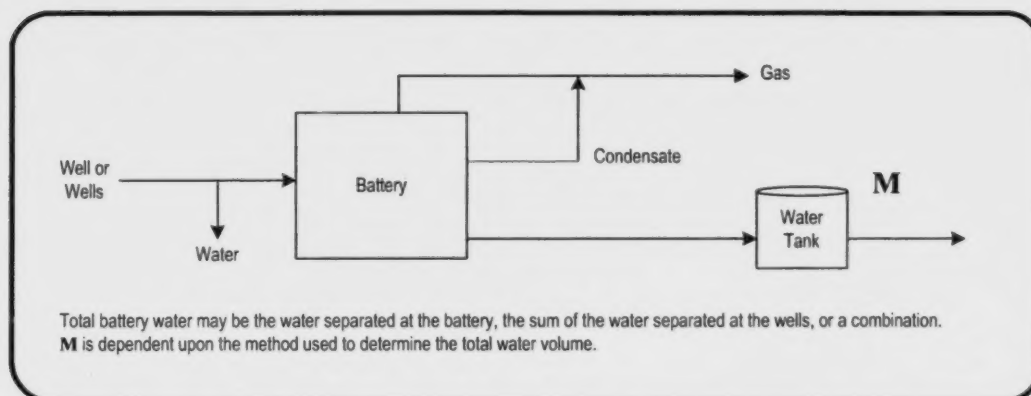
Figure 1.19

Maximum uncertainty of monthly volume = 5%

Single point measurement uncertainty = N/A
Total battery water may be determined by an individual group measurement, by totalling individual well measurements, or by totalling individual well estimates, so no basic requirement for measurement uncertainty has been set.

Total battery water in a gas system may be collected at a central location where it can be measured prior to disposal, or it may be a summation of individual well estimates or measurements of water collected at well sites and disposed from those sites. The 5% maximum uncertainty of monthly volume allows for some leeway in volume determination.

(xiii) Well water

For Figure 1.20,
(m) = single point measurement uncertainty
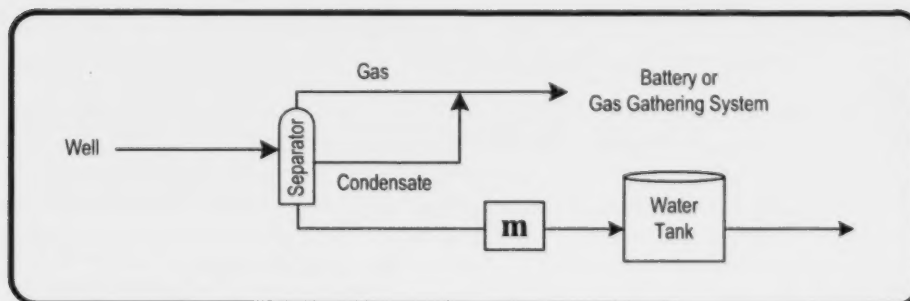
Figure 1.20

Single point measurement uncertainty = 10%

Maximum uncertainty of monthly volume = N/A
The uncertainty of the monthly volume will vary, depending upon whether produced volumes are subject to individual well measurement, estimation, or proration.

Water production at gas wells may be determined by measurement after separation, or if separators are not used, it may be determined by using water-gas ratios determined from engineering calculations or semiannual tests. To allow for the various methods used to determine production volumes, the single point measurement uncertainty is set at 10%.

1.7.3   Injection/Disposal Systems

(i) Total gas

For Figure 1.21,
M = maximum uncertainty of monthly volume
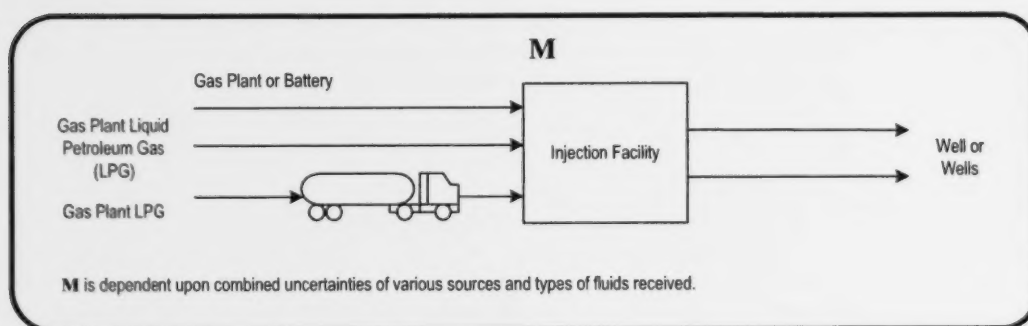
Figure 1.21

Maximum uncertainty of monthly volume = 5%

Single point measurement uncertainty = N/A
The single point measurement uncertainty will vary depending on the source and type of fluids received.

Gas used in injection/disposal systems may be clean processed gas or unprocessed gas that may contain entrained liquids, and in some cases several sources may make up the total gas volume received by an injection system. The expectation for the maximum uncertainty of monthly volume is set at 5% to allow for the uncertainties associated with measuring gas under those conditions.

(ii) Well gas

For Figure 1.22,
(m) = single point measurement uncertainty
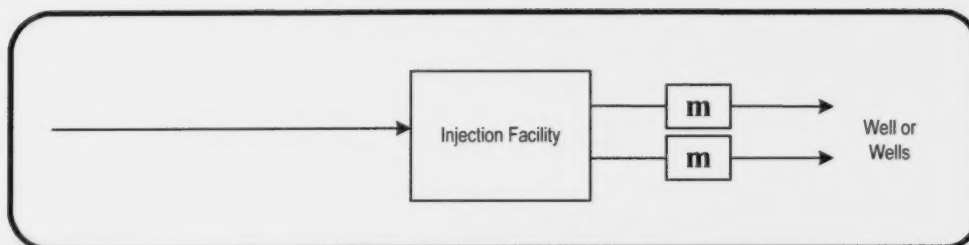
Figure 1.22

Single point measurement uncertainty = 3%

Maximum uncertainty of monthly volume = N/A
The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

The gas injected/disposed into wells must be measured and may consist of clean processed gas and/or unprocessed gas that may contain entrained liquids. The equipment and/or procedures used to determine the gas volumes injected/disposed into each well must be capable of meeting a 3% single point measurement uncertainty.

(iii) Total water

For Figure 1.23,
M = maximum uncertainty of monthly volume
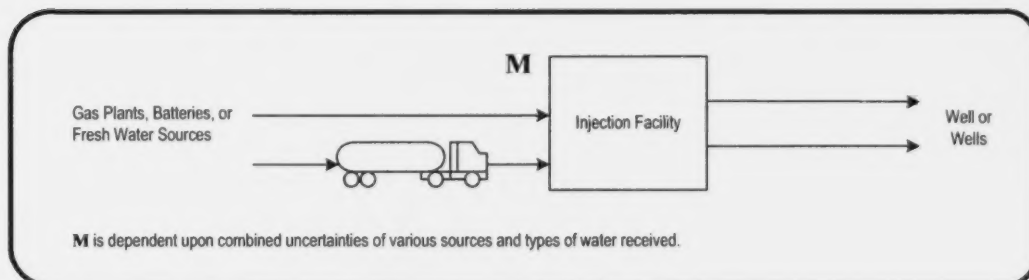
Figure 1.23

Maximum uncertainty of monthly volume = 5%

Single point measurement uncertainty = N/A
To be equivalent to the requirements for total oil and gas battery water.

Water used in injection/disposal systems may be produced water from oil or gas batteries, fresh water from water source wells, and/or waste water. To be equivalent to the requirements for total oil and gas battery water, the expectation for the maximum uncertainty of monthly volume is set at 5%.

(iv) Well water

For Figure 1.24,
(m) = single point measurement uncertainty
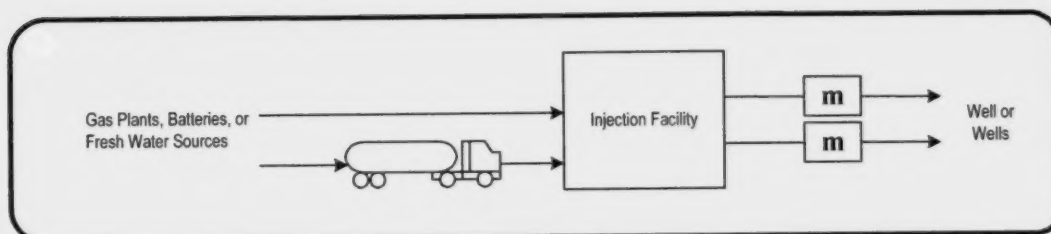
Figure 1.24

Single point measurement uncertainty = 5%

Maximum uncertainty of monthly volume = N/A
The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

The water injected/disposed into wells must be measured. The expectation for the single point measurement uncertainty is set at 5%.

1.8   Standards of Accuracy—Summary

1.8.1   Oil Systems

		Maximum uncertainty of monthly volume	Single point measurement uncertainty
(i)	Total battery oil (delivery point measurement)		
	Delivery point measures > 100 m^3/d	N/A	0.5%
	Delivery point measures ≤ 100 m^3/d	N/A	1%
(ii)	Total battery gas (includes produced gas that is vented, flared, or used as fuel)		
	> 16.9 10^3 m^3/d	5%	3%
	> 0.50 10^3 m^3/d but ≤ 16.9 10^3 m^3/d	10%	3%
	≤ 0.50 10^3 m^3/d	20%	10%
(iii)	Total battery water		
	> 50 m^3/month	5%	N/A
	≤ 50 m^3/month	20%	N/A
(iv)	Well oil (proration battery)		
	Class 1 (high), > 30 m^3/d	5%	2%
	Class 2 (medium), > 6 m^3/d but ≤ 30 m^3/d	10%	2%
	Class 3 (low), > 2 m^3/d but ≤ 6 m^3/d	20%	2%
	Class 4 (stripper), ≤ 2 m^3/d	40%	2%
(v)	Well gas (proration battery)		
	> 16.9 10^3 m^3/d	5%	3%
	> 0.50 10^3 m^3/d but ≤ 16.9 10^3 m^3/d	10%	3%
	≤ 0.50 10^3 m^3/d	20%	10%
(vi)	Well water	N/A	10%

1.8.2 Gas Systems

		Maximum uncertainty of monthly volume	Single point measurement uncertainty
(i)	Gas deliveries (sales gas)	N/A	2%
(ii)	Hydrocarbon liquid deliveries		
	Delivery point measures > 100 m^3/d	N/A	0.5%
	Delivery point measures ≤ 100 m^3/d	N/A	1%
(iii)	Plant inlet or total battery/group gas	5%	3%
(iv)	Plant inlet or total battery/group condensate (recombined)	N/A	2%
(v)	Fuel gas		
	> 0.50 10^3 m^3/d	5%	3%
	≤ 0.50 10^3 m^3/d	20%	10%
(vi)	Flare gas	20%	5%
(vii)	Acid gas	N/A	10%
(viii)	Dilution gas	5%	3%
(ix)	Well gas (well site separation)		
	> 16.9 10^3 m^3/d	5%	3%
	≤ 16.9 10^3 m^3/d	10%	3%
(x)	Well gas (proration battery)	15%	3%
(xi)	Well condensate (recombined)	N/A	2%
(xii)	Total battery water	5%	N/A
(xiii)	Well water	N/A	10%

1.8.3 Injection Systems

		Maximum uncertainty of monthly volume	Single point measurement uncertainty
(i)	Total gas	5%	N/A
(ii)	Well gas	N/A	3%
(iii)	Total water	5%	N/A
(iv)	Well water	N/A	5%

2 Calibration and Proving

2.1 Definitions

For the purposes of this directive, calibration and proving have distinctly different meanings:

Calibration refers to procedures or operations that establish the accuracy of the values indicated by a measuring device as compared to the values indicated by a calibration instrument that has accuracy traceable to a suitable measuring standard. Adjustments are made, if required, to the measuring device to make it comparable to the calibration instrument. Calibration may also refer to the procedure used to determine the volume of a prover.

Proving refers to the procedures or operations whereby a prover volume is compared to an indicated meter volume (both corrected to applicable pressure and temperature conditions). The prover volume divided by the indicated meter volume yields a meter factor. The meter factor is subsequently applied to indicate meter volumes to determine the adjusted volume. If the meter is connected to an electronic readout, the meter factor may be incorporated into the software such that the indicated meter volume is already corrected to read the adjusted volume. (Care should be taken in such cases not to apply the meter factor again. In such cases, the meter factor should be indicated on the tag or label as being 1.0000.)

A **meter factor** is a dimensionless number used to correct indicated meter volumes to adjusted volumes if the two volumes differ due to operational conditions (e.g., gas entrained in liquids, meter slippage, meter wear). It is not to be confused with the "K Factor," which is the factor used to convert the number of pulses generated by a meter whose operation is based on electronic pulse technology into units of volume. The K Factor is typically determined by the meter manufacturer and does not take into consideration all of the specific operational conditions the meter may be subjected to.

Prover refers to a device used to collect and determine the volume of a sample of fluid that has passed through a meter. Provers typically use volumetric or gravimetric means to determine the quantity of the sample.

A **prover run** refers to the operation of a prover or master meter whereby a representative volume is sampled and measured, and that sample volume is compared to the volume indicated by a meter through which the same sample has passed to determine a meter factor.

Master meter refers to a meter of known accuracy that is connected in series with another meter for the purpose of checking the accuracy of the second meter and providing a meter factor.

A **measuring standard** refers to a device used in calibration or proving that has a known value that is traceable to national reference standards maintained by the National Research Council in Ottawa, Ontario.

2.2 Applicability

The calibration and proving requirements stipulated in this directive are applicable to measurement devices used in Alberta to determine volumes for ERCB-required accounting and reporting purposes. These requirements are not applicable to measurement devices used only for a licensee's internal accounting purposes. The requirements are considered minimums, and a licensee may choose to apply more stringent requirements.

If a licensee wishes to deviate from these requirements or exceptions (other than applying more stringent requirements), see Section 1: Standards of Accuracy to determine if the deviation requires submission of an application to and approval by the ERCB.

2.3 Frequency

The accuracy of measurement devices may deviate over time, due to wear, changes in operating conditions, changes in ambient conditions, etc. Generally, the more important the accuracy of a measurement device is, the more frequently it must be calibrated or proved.

For the purposes of this directive, calibration or proving frequency have the following meanings:

- Calendar quarters are January to March, April to June, July to September, and October to December.
- Monthly means at least once per calendar month.
- Bimonthly means at least once every two calendar months.
- Quarterly means at least once per calendar quarter.
- Semiannually means at least once every two calendar quarters.
- Annually means at least once every four calendar quarters.
- Biannually means at least once every eight calendar quarters (once every two years).
- Triennially means at least once every twelve calendar quarters (once every three years).

Example: For an annual frequency, if the last calibration was performed in May 2006, the operator has to perform another calibration by the end of June 2007 (end of the calendar quarter).

Exceptions

1) If the use or operation of a measurement device requiring monthly or quarterly calibration/proving is suspended for a significant period (at least seven consecutive days), the scheduled calibration/proving may be delayed by the number of days the device was not in use. Documentation of the amount of time the device was not in service must be kept and made available to the ERCB on request. If this exception is being applied, the licensee must attach a tag to the meter indicating that this exception is in effect and the next scheduled calibration/proving date. This exception is not applicable to measurement devices subject to calibration/proving frequencies that are semiannual or longer.

2) If a liquid meter is removed from service for bench proving but is put "on the shelf" and not returned to service for a considerable period of time, the countdown to the next required bench proving does not start until the meter is returned to service. The licensee must attach a tag to the meter indicating the installation date, but leaving the original proving tag intact.

3) The ERCB may request that calibration/proving of a meter be done at any time or may extend the due date for scheduled calibration/proving, depending on the specific circumstances at a measurement point.

2.4   Accuracy of Provers and Calibration Instruments

Provers and other instruments used for calibration of measurement devices must be tested for accuracy prior to first being used, immediately following (prior to being used) any repairs or alterations being conducted on them, and periodically, in accordance with the following:

- Portable provers must be calibrated biannually using measurement standards that are traceable to national standards.

- Stationary provers must be calibrated every four years using measurement standards that are traceable to national standards.

- Calibration instruments, such as manometers, thermometers, pressure gauges, deadweight testers, electronic testers, etc., must be tested for accuracy biannually against instruments having accuracy traceable to national standards.

- Master meters must be proved quarterly using a calibrated prover. The fluid used to prove the master meter must have properties similar to the fluids measured by the meters it will be used to prove. The master meter must be proved at flow rates that are comparable to the conditions it will be used for.

The procedures to be followed for these accuracy tests must be designed to provide consistent and repeatable results and must take into consideration the actual operational conditions the device will encounter. To that end, the procedures must be in accordance with the following, as available and applicable (presented in order of ERCB preference from first to last):

- procedures specified by Measurement Canada (An Agency of Industry Canada),

- procedures described in the API *Manual of Petroleum Measurement Standards*,

- the device manufacturer's recommended procedures, or

- other applicable Industry-accepted procedures.

If none of the foregoing exists, the ERCB will consider applications for and may grant approval of, appropriate procedures.

Records of the foregoing accuracy tests must be kept for at least one year following the expiry of the applicable test and provided to the ERCB on request.

2.5   Gas Meters

2.5.1   General Calibration Requirements

The term "gas meter" is broadly used to describe all of the equipment or devices that are collectively used to arrive at an indication of a gas volume. Typically, various values (e.g., differential pressure, static pressure, temperature) must be determined and used to calculate a gas volume. Depending on the specific gas meter, each of those values may be determined by individual devices or equipment. In this section, "gas meter" means the main device through which gas flows and which typically provides an indication of flow. The term "meter element" means the devices that are associated with the gas meter but typically provide values other than the indication of flow. The term "end device" means the device or equipment that records the various values used to calculate a gas volume. In the case of electronic flow measurement, the end device may also perform the calculations necessary to arrive at the measured gas volume.

Typically, calibration of gas meter equipment requires the various meter elements to be subjected to various actual pressure, temperature, and other values that are concurrently subjected to the calibration equipment. If the end device does not indicate the same value as the calibration equipment, adjustments must be made to the meter element and/or end device.

Some meter equipment technologies may require alternative equipment and procedures for calibration, which is acceptable provided that the equipment and procedures are capable of confirming that the meter elements are functioning properly and are sensing and transmitting accurate data to the end device.

Orifice meters are commonly used to measure gas volumes in Alberta. Gas orifice meters themselves (the meter run and orifice plate-holding device) do not require calibration/proving. However, the associated meter elements and the end devices to which they are connected must be calibrated, as described in Section 2.5.3: Orifice Meters.

If devices other than orifice meters are used to measure gas, the associated meter elements and the end devices to which they are connected must be calibrated at the same frequency as orifice meters. The required procedures must be designed to provide consistent and repeatable results and must take into consideration the actual operational conditions the device will encounter. To that end, the procedures must be in accordance with the following, as available and applicable (presented in order of ERCB preference from first to last):

- procedures specified by Measurement Canada,
- procedures described in the API *Manual of Petroleum Measurement Standards* (*MPMS*),
- the device manufacturer's recommended procedures, or
- other applicable industry-accepted procedures.

If none of the foregoing exists, the ERCB will consider applications for and may grant approval of appropriate procedures.

The frequency of gas meter calibration must be

- within the first calendar month of operation of a new meter,
- immediately (by the end of the calendar month) following service or repairs to the meter,
- semiannually thereafter if the meter is used in a gas plant or for sales/delivery point (royalty trigger points), and
- annually for all other meters.

The royalty trigger points for gas are generally for clean processed gas sales at the plant gate or for raw gas that is sent to another facility for FUEL usage only.

2.5.2   Gas Meter Internal Inspection

A key contributor to meter accuracy is the condition of the internal components of the gas meter. Examples of internal components are orifice plates, vortex shedder bars, and turbine rotors. The internal components must be removed from service, inspected, replaced or repaired if found to be damaged, and then placed back in service, in accordance with the following:

- The required frequency for inspection of internal gas meter components is semiannually for gas plant accounting meters and sales/delivery point (royalty trigger point) meters and annually for all other gas meters.

- Whenever possible, the inspection should be done at the same time as the calibration of the meter elements and end device, but to accommodate operational constraints the inspection may be conducted at any time, provided that the frequency requirement is met.

- Inspections must be done in accordance with procedures specified by the API, the American Gas Association (AGA), or other relevant standards organizations, other applicable industry-accepted procedures, or the device manufacturer's recommended procedures, whichever are most applicable and appropriate.

- A tag or label must be attached to the meter or end device that identifies the meter serial number, the date of the internal inspection, and any other relevant details (e.g., the size of the orifice plate installed in the meter).

- A detailed record of the inspection documenting the condition of the internal components found and any repairs or changes made to the internal components must be kept for at least one year and provided to the ERCB on request.

2.5.2.1 Exceptions

1) If the "as found" calibration check of the instrumentation of a gas meter confirms that the accuracy of all readings or outputs are within ± 0.25% of full scale (with the exception of ± 1°C for the temperature element), no adjustment of the instrumentation is required.

2) If electronic gas meters have been found to not require adjustment for three consecutive calibrations, as indicated in #1 above, the minimum time between routine calibrations may be doubled. A tag must be attached to the meter indicating that this exception is being applied and the date of the next scheduled calibration. The records of the calibrations that qualify the meter for this exception must be kept for at least one year and made available to the ERCB on request.

3) If redundant gas meters are installed for a measurement point or redundant meter elements and/or end devices are installed on a single gas meter, the minimum time between routine calibration of the meter elements and end devices may be doubled, provided that daily volumes from each end device are compared at least monthly and found to be within 0.25% of each other. If the daily volumes are not found to be within 0.25% of each other, immediate calibration of both sets of equipment is required. A tag must be attached to the meter indicating that this exception is being applied and the date of the next scheduled calibration. The records of the monthly comparisons and any calibrations that are done must be kept for at least one year and made available to the ERCB on request.

4) If rotary or other types of gas meters with internal moving parts are used to measure gas, such as fuel gas, those meters must be proved at a frequency of once every seven years following an initial proving prior to installation. These meters must also be proved immediately following any repairs or alterations being conducted on them. The proving may be done with the meter in service, or the meter may be removed from service and proved in a shop at a pressure that is within the normal operating condition for that meter location unless it can be shown that proving at lower pressure conditions will not change the uncertainty of the meter, such as in the case of a rotary meter. A tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, and the meter factor determined by the proving. A detailed report indicating the details of the proving operation must be either left with the meter or readily available for inspection by

the ERCB. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)

5) For meters used in effluent (wet gas) measurement that require proving, such as a turbine meter, the proving must be performed by using a gas master meter or other provers in single-phase proving runs. For ECF-WGR testing, see Section 7.4.

6) If the internal components of gas meters have been found to be clean and undamaged for three consecutive inspections, the minimum time between inspections may be doubled. When the internal components are found to be dirty or damaged on any subsequent inspection, the frequency for inspections will revert back to the original requirement.

7) If the inspection of internal components of a gas meter requires the meter to be removed from service and there is no meter bypass installed, it is acceptable to defer a scheduled internal component inspection until the next time the gas meter run is shut down, provided that shutting down and depressuring the gas meter run to remove and inspect the internal components would be very disruptive to operations, require excessive flaring, or cause a safety concern, **and**

- historical internal component inspections have proven to be satisfactory, **or**
- the meter run is installed in a flow stream where the risk of internal component damage is low (e.g., sales gas, fuel gas), **or**
- the measurement system at the facility provides sufficient assurance, through volumetric and/or statistical analysis, that internal component damage will be detected in a timely manner.

8) In the case of an orifice meter, if the orifice plate is mounted in a quick-change (senior) orifice meter assembly and when attempting to conduct an inspection of the orifice plate the fitting is found to be leaking between the chambers such that the meter run must be shut down and depressured to safely remove the orifice plate, it is acceptable to defer a scheduled orifice plate inspection until the next time the gas meter run is shut down, provided that

- shutting down and depressuring the gas meter run to remove the orifice plate would be very disruptive to operations, require excessive flaring, or cause a safety concern, **and**
- the orifice meter assembly is scheduled for repairs to be conducted the next time the gas meter run is shut down to eliminate the cause of the leak and allow future scheduled orifice plate inspections to be conducted, **and**
  - historical orifice plate inspections have proven to be satisfactory, **or**
  - the meter run is installed in a flow stream where the risk of orifice plate damage is low (e.g., sales gas, fuel gas, etc.), **or**
  - the measurement system at the facility provides sufficient assurance, through volumetric and/or statistical analysis, that orifice plate damage will be detected in a timely manner.

9) Internal metering diagnostics may be used to determine if the primary measurement element is within acceptable operating parameters and checked at the same required intervals as an internal inspection. Then internal inspection is not required until an alarm or error is generated by the device or as recommended by the manufacturer. The operator

must maintain documentation on the diagnostic capability of the measurement system and make it available to the ERCB on request.

Note that should gas meter internal component inspections be deferred in accordance with any of the foregoing exceptions, the licensee must be able to demonstrate to the ERCB, on request, that the situation meets the conditions identified. If these exceptions are being used, that must be clearly indicated on a tag or label attached to the meter (or end device). Evidence in battery or facility logs that the internal component inspection has been scheduled for the next shutdown must be available for inspection by the ERCB. For the purposes of these exceptions, "shutdown" means any scheduled discontinuation of flow through the meter that is of sufficient duration to allow the operations needed to remove and inspect the internal component. If an unscheduled shutdown appears that it will allow sufficient time to conduct internal component inspection operations, the licensee should consider conducting those inspections prior to the conclusion of this unscheduled shutdown.

2.5.3 Orifice Meters

The procedure for orifice meter chart recorder (end device) calibration must be in accordance with the following:

- Pen arc, linkage, pressure stops, and spacing must be inspected and adjusted, if necessary.

- The differential pressure element must be calibrated at zero, full span, and nine ascending/descending points throughout its range. A zero check of the differential under normal operating pressure must be done before and after the calibration.

- The static pressure element must be calibrated at zero, 50% of full span, and full span.

- If a temperature element is in place, the temperature element must be calibrated at three points (operating temperature, one colder temperature, and one warmer temperature are recommended).

- If a thermometer is in place and used to determine flowing gas temperature, the thermometer must be checked at two points and replaced if found to not read accurately within ± 1°C (operating temperature and one other temperature are recommended).

- If a thermometer or other temperature measuring device is not left in place (transported by an operator and used to determine flowing gas temperatures at multiple sites), the accuracy of that device must be verified at the same frequency and in the same manner as a thermometer left in place, and the record of that verification must be readily available for inspection by the ERCB.

- Subsequent to the meter calibration, a tag or label must be attached to the meter (or end device) that identifies the meter serial number and the date of the calibration.

- A detailed report indicating the tests conducted on the meter during the calibration and the conditions as found and as left must be either left with the meter (or end device) or readily available for inspection by the ERCB. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)

2.5.4 EFM Calibration

The procedure for calibration of an electronic flow measurement system must be in accordance with the following:

- For digital output devices, the differential pressure element must be calibrated at zero, 50% of calibrated full span, and calibrated full span.

- For analog output devices, the differential pressure element must be calibrated at zero, 50% of full span, and full span (ascending), as well as 80% and 20% (or 75% and 25%) of full span (descending).

- A zero check of the differential under normal operating pressure must be done before and after the calibration.

- The static pressure element must be calibrated at zero, 50% of full span, and full span.

- If a temperature element is in place, the temperature element must be verified at two points (operating temperature and one colder or one warmer temperature) and calibrated or replaced if found to not indicate accurately within $\pm 1°C$.

- Subsequent to the meter calibration, a tag or label must be attached to the meter (or end device) that identifies the meter serial number and the date of the calibration.

- A detailed report indicating the tests conducted on the instruments during the calibration and the conditions as found and as left must be either left with the meter (or end device) or readily available for inspection by the ERCB. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)

- If data from the instrumentation are sent to another location(s) for flow calculations via DCS, SCADA, RTU, or other means of communication, the reading of the calibration must be verified at the receipt location of such data to ensure accurate data transmission.

2.6 Liquid Meters

Oil and other liquid production (except gas well condensate under certain conditions) and disposition volumes must always be reported as liquid volumes at 15°C and either equilibrium pressure (equilibrium pressure is assumed to be atmospheric pressure at the point of production or disposition) or 101.325 kilopascals (kPa) absolute pressure.

Meters used to measure hydrocarbons, water, and emulsions are subject to the following general proving requirements. However, there are additional specific requirements depending on the fluid types, as detailed in Sections 2.6 through 2.10.

1) The design and operation of the meter installation must ensure that the conditions of fluid flow through the meter are within the manufacturer's recommended operating range. The meter must be installed upstream of a snap acting control/dump valve, if present.

2) The size of the prover taps and operation of the prover must not restrict or alter the normal flow through the meter. Tank-type volumetric or gravimetric provers must be connected downstream of the meter and downstream of a snap acting control/dump valve, but other provers, such as ball provers, pipe provers, or master meters, may be connected either upstream (provided there is no gas breakout) or downstream of the meter before a snap acting control/dump valve.

3) A new hydrocarbon meter must be proved within the first calendar month of operation or immediately following any repairs being conducted on the meter or any changes to the meter installation. (Note that the resultant meter factor must be applied back to the volumes measured after the commencement of operation, repair, or change). A new water

meter must be proved within the first 3 months of operation or immediately following any repairs being conducted on the meter or any changes to the meter installation and no retroactive application of meter factor is required.

4) The new meter must be proved according to the frequency in Table 2.1 thereafter.

Table 2.1. Proving frequency requirements for hydrocarbons, water, and emulsions

Meter application	Fluid type	Proving frequency
Wellhead, group, injection	Test oil, test emulsion, live condensate, water	Annually
Gas plant, cross-border	Live condensate, NGL, LPG	Semiannually
Delivery point, LACT	Oil, condensate, live condensate, NGL, LPG	Monthly (see exceptions below)

5) The meter must be proved in line at normal operating conditions unless otherwise exempted by the ERCB.

6) If a master meter is used for proving, it must have an uncertainty rating equal to or better than the meter it is being used to prove.

7) Each proving run must consist of a representative volume of hydrocarbons or hydrocarbons/water emulsion being directed into the prover or through the master meter. After application of any required correction factors, the resultant volume determined by the prover or master meter is divided by the metered volume to determine the meter factor.

8) If a meter is proved after a period of regular operation, an "as found" proving run must be performed prior to conducting any repairs on the meter or replacing the meter.

9) An acceptable initial proving (the first proving of a new or repaired meter) and all subsequent proving must consist of the number of consecutive runs, each with a meter factor (MF) within the mean of all applicable runs, as specified in Table 2.1. The resultant meter factor will be the average of all the applicable run meter factors. (Proving procedures using more than the specified number of runs are allowed, provided that the licensee can demonstrate that the alternative procedures provide a meter factor of equal or better accuracy.)

Table 2.2. Proving requirements for hydrocarbons, water, and emulsions

Hydrocarbon meter type	Initial prove: number of required consecutive runs	Subsequent prove: number of required consecutive runs		Maximum MF deviation allowed from mean of all applicable runs (%)
		"As found" MF ≤ ± 0.5% of previous	"As found" MF >± 0.5% of previous	
Live oil – field proving (see 2.7)	4	1	4	1.5
Live oil – shop proving (see 2.7.1)	4	4	4	0.5
Dead oil, condensate at equilibrium, high vapour pressure liquids (see 2.7.2 & 2.8.1)	3	1	3	0.25
Live condensate – field proving (see 2.8.2)	4	1	4	2
Live condensate – shop proving (see 2.8.2.1)	4	4	4	0.5
Water – field proving (see 2.10)	4	1	4	1.5
Water – shop proving (see 2.10)	4	4	4	1.5

10) A detailed report indicating the type of prover or master meter used, the run details, and the calculations conducted during the proving must be either left with the meter or readily available for inspection by the ERCB. (If the detailed report is left with the meter, the requirement below relating to the tag or label is considered to be met.) If the proving involved the use of a shrinkage factor instead of degassing, a copy of the sample analysis must be attached to the proving report.

11) Subsequent to the meter proving, a tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, the type of prover or master meter used, and the average meter factor. If the meter is connected to an electronic readout, it may be possible to program the meter factor into the software to allow the meter to indicate corrected volumes. If the meter is connected to a manual readout, it is necessary to apply the meter factor to the observed meter readings to result in corrected volumes.

Exceptions

1) If a meter used to measure fluids at flow-line conditions is a type that uses no internal moving parts (e.g., orifice meter, vortex meter, V-cone meter), it does not require proving, provided that the following conditions are met:

- The flow through the meter must be continuous and maintained within the rates specified by the meter manufacturer as providing accurate measurement.

- The design and operation of the entire meter system must be in accordance with the meter manufacturer's specifications.

- The meter end device(s) must be calibrated at the frequencies specified in Section 2.5, using procedures specified in Section 2.5, by the API MPMS, the AGA, the device manufacturer, or other applicable industry-accepted procedures, whichever are most appropriate and applicable.

- The internal components of the primary meter device must be removed from service at the same frequency as indicated in Table 2.1, inspected, replaced or repaired if found to be damaged, and then placed back in service, in accordance with procedures specified by the API, the AGA, other relevant standards organizations, other applicable industry-accepted procedures, or the device manufacturer's recommended procedures, whichever are most applicable and appropriate. Internal metering diagnostics may be used to determine if the primary measurement element is within acceptable operating parameters and checked at the same required intervals as an internal inspection. Then internal inspection is not required until an alarm or error is generated by the device or as recommended by the manufacturer. The operator must maintain documentation on the diagnostic capability of the measurement system and make that available to the ERCB on request.

- If a meter is to be proved just like one with internal moving parts, no internal inspection is required.

- Whenever possible, the inspection of internal components should be done at the same time as the meter end device calibration, but to accommodate operational constraints the inspection may be conducted at any time, provided the frequency requirement is met.

- A tag or label must be attached to the meter (or end device) that identifies the primary device serial number and the date of the calibration.

- A tag or label must be attached to the meter (or end device) that identifies the primary device serial number, the date of the internal components inspection, and any other relevant details (e.g., the size of the orifice plate installed in the meter).

- A detailed report indicating the tests conducted on the meter during the calibration and the conditions as found and as left must be either left with the meter (or end device) or readily available for inspection by the ERCB. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)

- A detailed record of the internal components inspection documenting their condition as found and any repairs or changes made to them must be either left with the meter (or end device) or readily available for inspection by the ERCB. (If the detailed report is left with the meter or readily available, the foregoing requirement relating to the tag or label is considered to be met.)

2) If the volume of fluid measured by a delivery point or LACT meter does not exceed 100 m^3/d, the meter proving frequency may be extended to quarterly. The tag attached to the meter must clearly indicate that the meter measures ≤ 100 m^3/d and that the meter is on a quarterly proving frequency. The required proving frequency will revert back to monthly if the meter begins measuring volumes > 100 m^3/d.

3) For delivery point or LACT meters, if the meter factor has been found to be within 0.5% of the previous factor for three consecutive months, the meter proving frequency may be extended to quarterly. The tag attached to the meter must clearly indicate that the meter has been found to have consistent meter factors and is on a quarterly proving frequency. The required proving frequency will revert back to monthly whenever the meter factor determined during a proving is found to not be within 0.5% of the previous factor.

2.7   Oil Meters

The two basic ways that oil is measured require distinctly different proving procedures. If oil production is measured prior to being reduced to atmospheric pressure, the proving procedures must allow for the volume reduction that will occur when the gas in solution with the "live" oil is allowed to evolve upon pressure reduction. No consideration for gas in solution is required when proving meters used to measure "dead" oil.

2.7.1   Live Oil Meters

Live oil meters are typically those used to measure volumes of oil or oil/water emulsion produced through test separators, but they also include meters used to measure well or group oil or oil/water emulsions that are delivered to other batteries or facilities by pipeline prior to the pressure being reduced to atmospheric.

To account for the shrinkage that will occur at the metering point due to the gas held in solution with live oil, the proving equipment and procedures may determine the amount of shrinkage either by physically degassing the prover oil volumes or by calculating the shrinkage based on an analysis of a sample of the live oil. Calculation of shrinkage volumes is most often used to mitigate safety and environmental concerns if the live oil volumes are measured at high pressures or if the live oil contains hydrogen sulphide (H_2S).

Additional proving requirements for live oil are as follows.

1) If the proving procedure includes degassing the prover to physically reduce the pressure of the hydrocarbons to atmospheric pressure:

- The prover must be a tank-type volumetric or gravimetric prover.

- Each proving run must consist of a representative volume of hydrocarbons or hydrocarbons/water emulsion being directed through the meter and into the prover and the liquid volume then being reduced in pressure to atmospheric pressure. The resultant volume determined by the prover, after application of any required correction factors, is divided by the metered volume to determine the meter factor.

- The amount of time required to degas the prover volume and arrive at a stable atmospheric pressure in the prover will vary, depending on the initial fluid pressure and the fluid characteristics.

2) If the proving procedure uses a shrinkage factor, rather than degassing, to adjust the prover volume to atmospheric conditions:

- A shrinkage factor representative of the fluid passing through the meter must be determined and used to adjust the meter volumes to atmospheric conditions.

  - The shrinkage factor may either be incorporated into the meter factor or be applied to metered volumes after they are adjusted by the meter factor.

  - The shrinkage factor must be based upon analysis of a sample of the metered fluid taken at normal operating conditions (see Section 6.3.2.3).

- Whenever the operating conditions at the meter experience a change that could significantly affect the shrinkage factor, a new shrinkage factor must be determined based upon analysis of a sample of the metered fluid taken at the new operating conditions. Consideration should also be given to proving the meter at the new operating conditions to determine if the meter factor has been affected.

- When the shrinkage factor option is used, the tag attached to the meter must indicate that a shrinkage factor was used instead of degassing the prover and whether the shrinkage factor was incorporated into the meter factor or will be applied separately.

3) When proving a test oil meter, a well that is representative of the battery's average well production characteristics must be directed through the test separator for each of the four runs. If there are wells in the battery with production characteristics that vary significantly from the average, consider determining specific meter factors to be used for each of those wells.

4) In the case of a test oil meter, the meter factor must include a correction factor to adjust the metered volume to 15°C (unless the meter is temperature compensated). Although the actual fluid temperature may vary with ambient temperature, it is acceptable to assume that the temperature observed at the time of proving is reasonably representative of the temperature experienced at the meter until the next proving. This requirement does not apply to meter technologies that do not require correction for temperature.

5) In the case of a live oil delivery point meter, the meter factor must not include a correction factor for temperature. The meter must either be temperature compensated or a fluid temperature must be taken daily and the metered volume must be corrected to 15°C. This requirement does not apply to meter technologies that do not require correction for temperature.

Exceptions

1) In situations where individual well production rates are so low that proving a test oil meter in accordance with the requirements listed above would require excessive time, it is acceptable to modify the proving procedures. Complete, individual proving runs

requiring more than one hour are considered excessive. The following modifications, in order of ERCB preference, may be used to reduce proving time:

- Produce several wells through the test separator at one time to increase the volume available for the proving runs.

- If the degassing procedure is being used, degas the first run only, and then use those data to calculate a shrinkage factor, which can be applied to subsequent runs conducted without degassing.

- Use the highest rate well for all proving runs.

- Conduct only three proving runs.

Note that the detailed proving report must clearly indicate if any of the foregoing modifications was used to prove the meter.

2) A live oil meter may be removed from service and proved in a meter shop, if it meets the following:

- If the meter is used to measure test volumes of conventional oil/emulsion, the average rate of flow of oil/emulsion of all the wells tested through the meter must be ≤ 2 m^3/d and no well may exceed 4 m^3/d of oil/emulsion production, or

- any meter used to measure test volumes of heavy oil/emulsion (density > 920 kg/m^3) or crude bitumen may be proved in a meter shop.

3) Shop proving is to be conducted in accordance with the following in addition to the general procedure in Section 2.6 where applicable:

- The meter installation must be inspected as follows, and corrective action must be taken when required:

  - The flow rate through the meter must be observed to verify that it is within the manufacturer's recommended operating ranges.

  - The dump valve must not be leaking (no flow registered between dumps).

- The shop proving may be conducted with a volumetric or gravimetric prover or with a master meter, as follows:

  - Water is typically used as the proving fluid, but varsol or some other light hydrocarbon fluid may be used for the proving.

  - Corrections for the temperature and pressure of the proving fluid must be made, where applicable.

If the gas held in solution with the fluid produced through the meter is of sufficient volume to significantly affect the fluid volume indicated by the meter, consideration should be given to determining an appropriate shrinkage factor to correct for the effect of the solution gas and provide that factor to the meter calibration shop so it may be built into the meter factor.

4) If a meter that required internal inspection is used to measure liquid hydrocarbon, the exception under item 7 in 2.5.2.1 also applies.

2.7.2 Dead Oil Meters

Dead oil meters are typically those used for delivery point (or custody transfer point) measurement of clean oil that has been degassed to atmospheric pressure. These meters may be found measuring oil being pumped from a battery into a pipeline or measuring oil being pumped from a truck into a pipeline terminal, battery, or other facility.

2.8 Condensate Meters

Condensate is subject to two different sets of measurement, accounting, and reporting rules. If condensate volumes are measured and delivered at atmospheric pressure or equilibrium pressure, the volume must be determined and reported as a liquid volume at 15°C and equilibrium pressure. (Equilibrium pressure is assumed to be either atmospheric pressure at the point of production or disposition or the actual equilibrium pressure.) If condensate volumes are measured and delivered at flow-line conditions, the volume must be determined at flow-line pressure and corrected to 15°C, but the volume is reported as a gas equivalent volume at standard conditions (101.325 kPa absolute and 15°C).

2.8.1 Condensate at Equilibrium Conditions

Meters that measure condensate stored and delivered as a liquid at atmospheric pressure or equilibrium pressure are typically "delivery point" meters and are therefore subject to the same proving requirements and exceptions applicable to meters used for dead oil measurement (see Sections 2.6 and 2.7.2).

2.8.2 Condensate at Flow-line Conditions

When a meter that requires proving is used to measure condensate at flow-line conditions, it must be subjected to the proving requirements in Section 2.6.

2.8.2.1 Exceptions

A meter used to measure condensate at flow-line conditions may be removed from service and proved in a meter shop, in accordance with the following:

- If the meter is used to measure condensate production on a continuous basis, the rate of flow through the meter must be ≤ 2 m³/d or it must be ≤ 3 m³/d with the gas equivalent volume of the daily condensate volume being $\leq 3\%$ of the daily gas volume related to the condensate production. If the meter is used on a portable test unit, there is no volume limitation, but consideration should be given to proving the meter in line if significant condensate production is observed during the test.

- The meter installation must be inspected as follows, and corrective action must be taken where required:
  - The flow rate through the meter must be observed to verify that it is within the manufacturer's recommended operating ranges.
  - The dump valve must not be leaking (no flow registered between dumps).

2.9 Other Liquid Hydrocarbon Meters

Meters used to measure other high vapour pressure liquid hydrocarbons, such as propane, butane, pentanes plus, natural gas liquid/liquid petroleum gas (NGL/LPG), etc., are subject to the same proving requirements and exceptions as are meters used for measurement of condensate at equilibrium conditions or measurement of dead oil (see Section 2.6).

2.10 Water Meters

If a meter is used to measure water production, injection, or disposal or injection of other water-based fluids, in addition to the requirements in Section 2.6,

- the meter must be installed and proved within the first three months of operation (note that the meter factor may be assumed to be 1.0000 until the first proving is conducted); and

- the proving may be conducted in line at field operating conditions, or the meter may be removed from service and proved in a meter shop using water as the test fluid.

The proving may be conducted using a volumetric prover, a gravimetric prover, or a master meter.

If a meter is proved after a period of regular operation, an "as found" proving run must be performed prior to conducting any repairs on the meter or replacing the meter. An acceptable proving must consist of four consecutive runs (one of which may be the "as found" run), each providing a meter factor within ± 1.5% of the mean of the four factors. The resultant meter factor is the average of the four applicable meter factors. (Proving procedures using more than four runs will be allowed, provided that the licensee can demonstrate that the alternative procedures provide a meter factor of equal or better accuracy.)

Following the meter proving, a tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, and the average meter factor. If the meter is connected to an electronic readout, it may be possible to program the meter factor into the software to allow the meter to indicate corrected volumes. If the meter is connected to a manual readout, it is necessary to apply the meter factor to the observed meter readings to result in corrected volumes.

A detailed report indicating the type of prover or master meter used, the run details, and the calculations conducted during the proving must be either left with the meter or readily available for inspection by the ERCB. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)

2.11 Product Analyzers

If a product analyzer (water cut analyzer) is used to determine water production, it must be calibrated annually using procedures recommended by the manufacturer.

Following the calibration, a tag or label must be attached to the product analyzer that identifies the analyzer serial number and the date of the calibration. A detailed report indicating the calibration procedure used and the calibration details must be either left with the analyzer or readily available for inspection by the ERCB. (If the detailed report is left with the analyzer or readily available, the foregoing requirement relating to the tag or label is considered to be met.)

2.12 Automatic Tank Gauges

2.12.1 Inventory Measurement

If automatic tank gauge devices are used to indicate fluid levels in tanks for monthly inventory measurement, they must be calibrated on site within the first month of operation. The calibration procedures must be in accordance with the following, as available and applicable (presented in order of ERCB preference from first to last):

- the device manufacturer's recommended procedures,

- procedures described in the API *Manual of Petroleum Measurement Standards*, or

- other applicable industry-accepted procedures.

If none of the foregoing exists, the ERCB will consider applications for and may grant approval of appropriate procedures.

A record of the calibration must be made available to the ERCB on request.

2.12.2 Delivery Point Measurement

If automatic tank gauge devices are used to indicate fluid levels in tanks for delivery point measurement of oil or oil/water emulsion, such as truck volume receipts at batteries/facilities or batch deliveries into a pipeline, they must be calibrated on site within the first month of operation and monthly thereafter. The calibration procedures must be in accordance with the following, as available and applicable (presented in order of ERCB preference from first to last):

- the device manufacturer's recommended procedures,

- procedures described in the API Manual of Petroleum Measurement Standards, or

- other applicable industry-accepted procedures.

If none of the foregoing exists, the ERCB will consider applications for and may grant approval of appropriate procedures.

A record of the calibration must be made available to the ERCB on request.

Exception

Where the accuracy of an automatic tank gauge is found to be within 0.5% of full scale for three consecutive months, the calibration frequency may be extended to quarterly. The record of calibration must clearly indicate that the device has been found to demonstrate consistent accuracy and is on a quarterly calibration frequency. The records of the calibrations that qualify the device for this exception must be kept and made available to the ERCB on request. The calibration frequency will revert back to monthly whenever the accuracy is found to not be within 0.5% of full scale.

2.13 Using Tank Gauging for Oil Measurement

Tank gauging refers to determining levels in a tank and using those levels to calculate a volume increase or decrease in the tank. The level may be determined by using an automatic tank gauge device or by manually determining the level with a gauge tape. In either case, the volume of the tank relative to its height at any given point must be determined. This is referred to as the tank calibration, or tank strapping, and results in the creation of a tank gauge table.

2.13.1 Inventory Measurement

If tank gauging is used only for monthly inventory measurement, specific tank calibration procedures are not required. It is acceptable to use gauge tables provided by the tank manufacturer or, if those are unavailable, generic gauge tables applicable to the tank size/type being used.

2.13.2 Delivery Point Measurement

If tank gauging is used for delivery point measurement of oil or oil/water emulsion, such as truck volume receipts at batteries/facilities or batch deliveries into a pipeline, the specific tanks being used must be calibrated on site within the first month of operation and any time the tank is damaged or altered. The calibration must result in the creation of a gauge table for each tank, which must then be used in conjunction with tank gauge readings to determine volumes. Calibration procedures must be in accordance with applicable methods stipulated in the API *Manual of Petroleum Measurement Standards*.

A record of the calibration must be made available to the ERCB on request.

2.14 Weigh Scales

Weigh scales used to measure oil/water emulsion and clean oil receipts at batteries, custom treating plants, pipeline terminals, and other facilities must be approved and inspected prior to use, in accordance with Measurement Canada requirements.

Weigh scales must be tested for accuracy in accordance with the following schedule:

- monthly,
- immediately (by the end of the calendar month) following any incident in which the scale may have been damaged, and
- immediately (by the end of the calendar month) following any changes or modifications being made to the scale.
- The complete set of procedures set out by Measurement Canada for determining weigh scale accuracy must be used following any damage or modifications and at least annually.

The monthly accuracy tests may be done using the complete set of procedures set out by Measurement Canada or, as a minimum, using the following abbreviated procedure:

1) Zero check: Determine if the scale reads zero with no weight on the scale.

2) Add a 10 kg standard weight: Determine if the scale reads 10 kg.

3) Remove the 10 kg standard weight: Determine if the scale returns to zero.

4) Add a test load consisting of 10 000 kg of standard weights or, alternatively, durable object(s) of known weight (minimum 5000 kg): Determine if the scale reads the correct weight of the test load (acceptable error is ± 0.2% of the test load).

5) Add a loaded truck, typical of the loads routinely handled by the scale: Note the total weight of the test load and truck.

6) Remove the test load and note the weight of the truck alone: Determine if the scale reading correctly indicates the removal of the test load (acceptable error is ± 0.2% of the test load).

7) Remove the truck: Determine if the scale returns to zero with no weight on the scale.

If as a result of the foregoing tests the weigh scale is found to not be accurate, it must be calibrated and retested until found to be accurate and then sealed by a heavy-duty scale service company. The service company must then send a written report to Measurement Canada documenting the adjustment and/or repairs.

A detailed record of the accuracy tests and any calibration activities must be kept in close proximity to the weigh scale, retained for at least one year, and made available to the ERCB on request. This record must include the following information:

- make, model, serial number, and capacity of the weigh scale and any associated equipment,
- date of the accuracy test,
- details of the tests performed and the results noted, and
- details regarding any alterations or calibration performed on the weigh scale.

Exceptions

1) If the volume of fluid measured by a weigh scale does not exceed 100 m^3/d, the monthly accuracy test frequency may be extended to quarterly. The detailed record of the accuracy tests must clearly indicate that the weigh scale measures ≤ 100 m^3/d and that the weigh scale is on a quarterly testing frequency. The required testing frequency will revert back to monthly if the weigh scale begins measuring volumes in excess of 100 m^3/d.

2) If the weigh scale has been found to not require calibration adjustments for three consecutive months, the monthly accuracy test frequency may be extended to quarterly. The required accuracy test frequency will revert back to monthly whenever a quarterly accuracy test determines that the weigh scale requires calibration adjustments.

3 Proration Factors, Allocation Factors, and Metering Difference

3.1 Proration Factors and Allocation Factors

3.1.1 Description

Proration is an accounting system or procedure where the total actual monthly battery production is equitably distributed among wells in the battery. This system is applicable when the production of wells producing to a battery is commingled before separation and measurement, and each well's monthly production is initially estimated, based on well test data. In this type of system, proration factors are used to correct estimated volumes to actual volumes.

In the case of an oil proration battery (Figure 3.1), the oil, gas, and water produced by individual wells are not continuously measured. Instead, the wells are periodically tested to determine the production rates of oil, gas, and water. The rates determined during the well test are used to estimate the well's production for the time period beginning with the well test and continuing until another test is conducted. The estimated monthly production so determined for each well in the battery is totalled to arrive at the battery's total monthly estimated production. The total actual oil, gas, and water production volumes for the battery are determined, and for each fluid, the total actual volume is divided by the total estimated production to yield a "proration factor." The proration factor is multiplied by each well's estimated production to yield the well's actual production. Similar accounting procedures are used for gas batteries subject to proration.

Test rates are used to estimate monthly well production volumes of each product.

Estimated monthly battery production of each product is determined by totalling all wells' estimated production.

Actual monthly battery production volume of each product is determined by measured delivery and inventory changes.

For each product,
Proration Factor = Actual Battery Production / Estimated Battery Production

For each product for each well,
Actual Monthly Well Production = Estimated Monthly Well Production x Proration Factor

$\mathbf{m}$ = Measurement Point

Figure 3.1

An "allocation factor" is a type of proration factor. It is used at facilities where only fluids received by truck are handled, such as custom treating plants and third-party-operated clean oil terminals (Figure 3.2). The name of the factor has been chosen to reflect the differences between batteries that receive fluids from wells through flow lines (where proration factors are used) and facilities that receive fluids from batteries only by truck (where allocation factors are used). The purpose of an allocation factor is similar to a proration factor, in that it is used to correct fluid receipt volumes (considered estimates) to actual volumes based on disposition measurements taken at the outlet of the facility (and also considering inventory change). The allocation factor is determined by dividing the monthly total actual volume for each fluid by the monthly total estimated volume for each fluid. The total estimated volume of each fluid received from each source is multiplied by the allocation factor for that fluid to yield the actual volume received from that source.

Allocation Factor

Receipts | Dispositions

Custom Treater or Third-Party Clean Oil Terminal

Oil → m → Pipeline

Water → m → Injection/Disposal

Estimated receipts is determined by measuring each truckload of fluid.

Actual disposition is determined by measuring each product delivered and inventory changes.

For each product,
Allocation Factor = Actual Disposition / Estimated Receipt

For each product,
Actual Monthly Receipt from Each Facility = Total Estimated Monthly Receipt from Each Facility x Allocation Factor

m = Measurement Point

Figure 3.2

The allocation factors discussed in this section are not to be confused with the process whereby products delivered out of a gas plant are "allocated" back to each well in the system, based on individual well production volumes and gas analyses.

Measurement accuracy and uncertainty generally relate to random errors and, as such, are not directly comparable to proration and allocation factors, which generally relate to bias errors. The ERCB Standards of Accuracy (Section 1) focus on specific measurement points (i.e., inlet or outlet), whereas proration and allocation factors relate to a comparison of inlet (or estimated production) to outlet measurement. It is important to note that target factors for different products may be different due to the products being subjected to different levels of uncertainty. For example, the target factors for oil and water in a conventional oil proration battery are different, because while the estimated production volumes of oil and water are determined by the same type of measurement, the outlet volumes of the clean oil and water are not determined by the same type of measurement.

When measurement equipment and procedures conform to all applicable standards, it is assumed that the errors that occur in a series of measurements will be either plus or minus

and will cancel each other out to some degree. Where a bias error occurs in a series of measurements, there will be no plus/minus and all of the measurements are assumed to be in error by the same amount and in the same direction. Proration factors and allocation factors are therefore used to equitably correct all measurements for biased errors.

3.1.2 Target Factors

If measurement and accounting procedures meet applicable requirements, any proration factor or allocation factor should be acceptable, since it is assumed that the factor will correct for a bias error that has occurred. However, the ERCB expects proration factors and allocation factors to be monitored by operators and used as a "warning flag" to identify when the measurement system at a battery or facility is experiencing problems that need investigation.

The ERCB deems the ranges of proration factors and allocation factors indicated below to be acceptable targets. When a factor is found to exceed these limits, the licensee is expected to investigate the cause of the factor being outside the target range and document the results of the investigation and the actions taken to correct the situation. The ERCB acknowledges that in some batteries or facilities, physical limitations and/or the economics applicable to a particular situation may prohibit the resolution of situations where factors are consistently in excess of the targets indicated below. In that case, the licensee must also document the reason(s) that prohibit further action from being taken. This information does not have to be routinely submitted to the ERCB, but must be available to the ERCB on request for audit.

If the cause of a factor being outside these ranges is determined and the error can be quantified, the ERCB expects the reported production data to be amended, thereby bringing the factor back into line. If the cause is determined and action is taken to correct the situation for future months, but the findings are not quantifiable for past months, no amendments need to be submitted.

3.1.3 Exception

An exception to the foregoing procedure is allowed for conventional oil proration batteries if based on average rates determined semiannually,

- all wells in the battery produce ≤ 2 m^3/d of oil, or

- the majority of the wells in the battery produce ≤ 2 m^3/d of oil and no well produces > 6 m^3/d of oil.

In this case the licensee should still be aware of the proration factors and take corrective action where necessary, but need not expend a great deal of effort to conduct an investigation and document the results.

3.1.4 Acceptable Proration Factors and Allocation Factor Ranges

Proration Factors

Conventional oil battery
- oil = 0.95000 – 1.05000
- gas = 0.90000 – 1.10000
- water = 0.90000 – 1.10000

Heavy oil battery – primary production and waterflood operations
- oil = 0.85000 – 1.15000
- water = 0.85000 – 1.15000
- gas = no stated expectation due to generally low production volumes

Heavy oil battery – thermal recovery operations
- oil = 0.75000 – 1.25000
- water = 0.75000 – 1.25000
- gas = no stated expectation due to the nature of thermal production

Gas battery – dry gas production
- gas = 0.90000 – 1.10000

Gas battery – effluent measurement
- gas = 0.90000 – 1.10000
- water = 0.90000 – 1.10000

Allocation Factors

Custom treating plant
- oil = 0.95000 – 1.05000
- water = 0.90000 – 1.10000

Clean oil terminal (third-party operated, where applicable)
- oil = 0.95000 – 1.05000

3.2 Metering Difference

3.2.1 Description

For ERCB production reporting purposes, a "metering difference" is used to balance, on a monthly basis, any difference that occurs between the measured inlet/receipt volumes and the measured outlet/disposition volumes at a facility. Metering difference is generally acceptable as an accounting/reporting entity if a difference results from two or more measurements of the same product. Metering differences occur because no two measurement devices provide the exact same volume, due to the uncertainties associated with the devices. However, a more significant cause of metering differences is that the product measured at the inlet to a facility is usually altered by the process within the facility, resulting in a different product or products being measured at the outlet of the facility. It should be noted that metering difference differs from proration and allocation factors in that for batteries or facilities where those factors are used, the difference occurs between "estimated" and "actual" volumes.

Metering difference may be used as follows:

Injection/disposal systems (Figure 3.3)—Receipts into these facilities are typically measured prior to being split up and delivered to individual wells, where each well's volume is measured prior to injection/disposal.
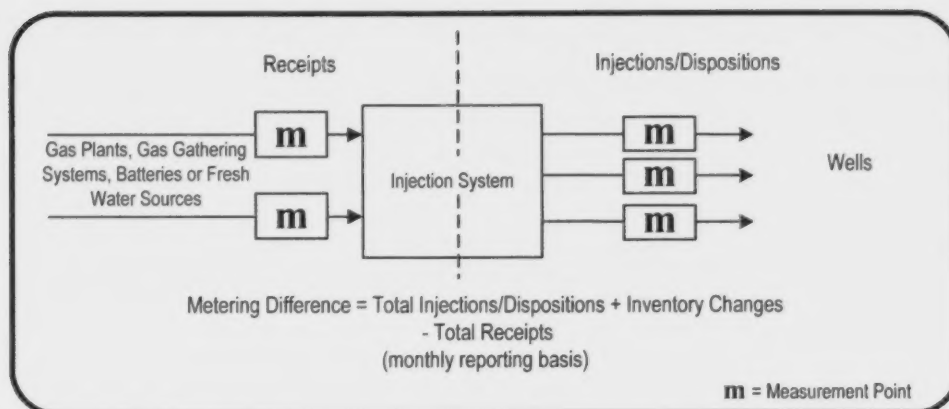
Figure 3.3

Batteries (Figure 3.4)—Metering difference may be used for gas and water production only, and only in limited, specific situations where there is both inlet and outlet measurement for reporting purposes. Such an example is a crude oil group battery where each well's gas production is measured and the combined gas stream is measured again before being sent to a gas gathering system or gas plant. However, metering difference would rarely be appropriate for use in a proration battery.

Gas gathering systems (Figure 3.4)—Receipts into these facilities are typically measured prior to being subjected to some sort of limited processing, which may include liquids removal and compression, and the resultant product(s) is measured prior to delivery to a sales point or to a gas plant for further processing.

Gas plants (Figure 3.4)—Receipts into these facilities are typically measured prior to being processed into salable products, and those products are measured prior to delivery to a sales point.
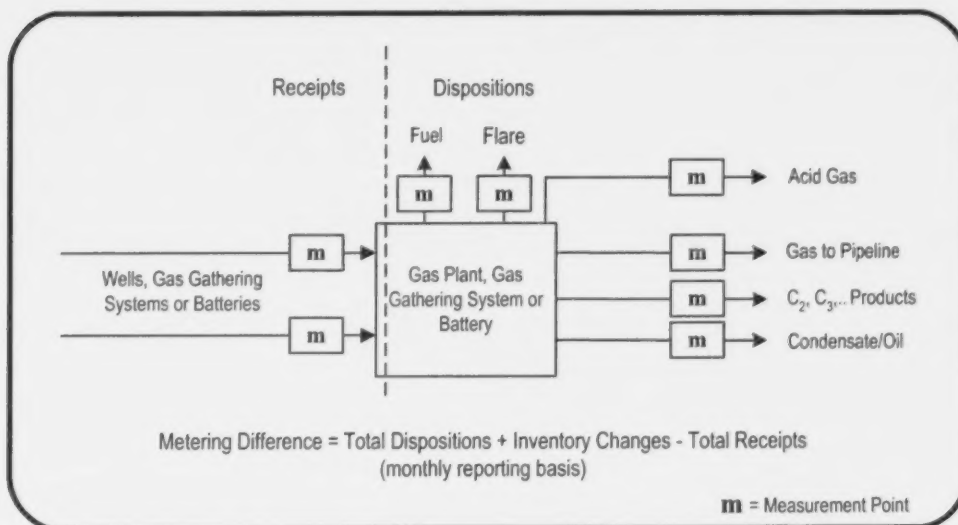
Figure 3.4

3.2.2   Target Metering Difference

If measurement and accounting procedures meet applicable requirements, metering differences up to ± 5% of the total inlet/receipt volume are deemed to be acceptable. The ERCB expects the metering difference to be monitored by licensees and used as a warning flag to identify when the measurement system at a battery or facility is experiencing problems that need investigation.

When a metering difference is found to exceed 5%, the licensee is expected to investigate the cause of the poor metering difference and document the results of the investigation and the actions taken to correct the situation. The ERCB acknowledges that in some batteries or facilities, physical limitations and/or the economics applicable to a particular situation may prohibit the resolution of situations where the metering difference is consistently in excess of the target indicated. In such cases, the licensee must also document the reason(s) that prohibit further action from being taken. This information does not have to be routinely submitted to the ERCB, but must be available to the ERCB on request for audit.

If the cause of a poor metering difference is determined and the error can be quantified, the ERCB expects the incorrectly reported production data to be amended, thereby bringing the metering difference back into line. If the cause is determined and action is taken to correct the situation for future months, but the findings are not quantifiable for past months, no amendments need to be submitted.

4 Gas Measurement

This section presents the base requirements and exceptions for gas measurement from any source in the upstream and midstream oil and gas industry that are used for determining volumes for reporting to the ERCB.

4.1   General Requirements

All gas production and injection must be continuously and accurately measured with a measurement device or determined by engineering estimation if exception conditions described below are met or site-specific ERCB approval has been obtained.

A gas measurement system is in compliance if the following base requirements are met. A licensee may deviate from these base requirements if the conditions in Section 4.3.5: Exceptions are met. Section 1: Standards of Accuracy provides further information relating to situations when other deviations may be acceptable.

Monthly gas volumes must be reported in units of 10^3 m^3 and rounded to 1 decimal place.

Standard or base conditions for use in calculating and reporting gas volumes are 101.325 kPa (absolute) and 15°C.

4.2   Gas Measurement and Accounting Requirements for Various Battery/Facility Types

4.2.1   Oil/Bitumen Batteries

4.2.1.1 General

- All wells in the battery must be classified as oil or bitumen wells.

- All wells in a multiwell battery must be subject to the same type of measurement. If there is a mixture of measured and prorated wells within the same battery, ERCB exception criteria in Section 5: Site-specific Deviation from Base Requirements under "Measurement by Difference" must be met or ERCB site-specific approval must be obtained.

- Production from gas batteries or other oil/bitumen batteries may not be connected to an oil/bitumen proration battery upstream of the oil/bitumen battery group gas measurement point unless specific criteria are met and/or ERCB approval of an application is obtained. See Section 5: Site-specific Deviation from Base Requirements under "Measurement by Difference" for examples.

Single-well Battery
- Gas must be separated from oil/bitumen or emulsion and measured (or estimated where appropriate) as a single phase.

Multiwell Group Battery
- Each well must have its own separation and measurement equipment, similar to a single-well battery.

- All equipment for the wells in the battery must share a common surface location.

Proration Battery
- All well production is commingled prior to the total battery gas being separated from oil/bitumen or emulsion and measured (or estimated where appropriate) as a single phase.

- Individual monthly well gas production is estimated based on periodic well tests and corrected to the actual monthly volume through the use of a proration factor.

4.2.2   Gas Batteries/Facilities

General

- All wells in the battery must be classified as gas wells.

- Gas wells may produce condensate or oil.

- All wells in a multiwell battery must be subject to the same type of measurement. If there are mixtures of measured and prorated wells (mixed measurement) within the same battery, ERCB exception criteria in Section 5: Site-specific Deviation from Base Requirements under "Measurement by Difference" must be met, or ERCB site-specific approval must be obtained, and the measured well(s) must have their own separate battery code(s) to deliver gas into the proration battery. Conversely, well(s) with no phase-separated measurement, including effluent wells, is not allowed to tie into a multiwell group battery unless there is a group measurement point before the tie-in.

- All wells in a multiwell battery must be connected by pipeline to a common point.

- Gas production from oil/bitumen wells or batteries or from other gas wells or batteries may not be connected to a gas proration battery upstream of the gas proration battery group measurement point unless ERCB exception criteria in Section 5: Site-specific Deviation from Base Requirements under "Measurement by Difference" are met or ERCB site-specific approval is obtained.

- Gas wells that are designed to operate on an on/off cycle basis using plunger lifts, on/off controllers, manual on/off, etc., or pumpjacks must report well status on the PRA as GAS PUMP instead of GAS FLOW.

- Any oil and gas facility, such as a well site, gas plant, battery, or individual compressor site, that is **designed** to consume fuel gas exceeding $0.5 \ 10^3 \ m^3/d$ must have fuel gas measurement from May 7, 2007, onwards. If it is a part of another facility on the same site, the overall site fuel gas used must be measured. The $0.5 \ 10^3 \ m^3/d$ measurement limit also applies to flare and vent gas streams, excluding heavy oil and crude bitumen batteries.

Single-well Battery

- Gas must be separated from water and condensate or oil (if applicable) and continuously measured as a single phase.

- Condensate produced must be reported as a liquid if it is disposed of from the well site without further processing.

- Condensate that is recombined with the gas production after separation and measurement or trucked from the well site to a gas plant for further processing must be converted to a gas equivalent volume and added to the measured single-phase gas volume for reporting purposes.

- Oil produced in conjunction with the gas must be reported as oil at stock tank conditions. The gas-in-solution (GIS) with the oil at the point of measurement must be estimated and added to the gas production volume. See Section 4.3.5.4.

Multiwell Group Battery

- Each well must have its own separation and measurement equipment, similar to a single-well battery.

- The wells in the group battery may all be identical with regard to handling of condensate, or there may be a mixture of methods for handling condensate. The rules for reporting condensate as a gas equivalent or as a liquid are the same as those for single-well gas batteries (above).

Multiwell Effluent Proration Battery

- The production from each well is subject to total effluent (wet gas) measurement, without separation of phases prior to measurement.

- Estimated well gas production is the effluent metered volume multiplied by an effluent correction factor (ECF) that is determined from periodic tests conducted at each well whereby a test separator is connected downstream of the effluent meter and the volumes measured by the test separator are compared to the volume measured by the effluent meter.

- Estimated well water production is determined by multiplying the water gas ratio (WGR), which is determined from the periodic tests, by the estimated well gas production.

- The combined (group) production of all wells in the effluent proration battery must have three-phase separation or equivalent and be measured as single-phase components, and the resulting total actual battery gas volume (including gas equivalent volume [GEV] of condensate) and total actual battery water volume must be prorated back to the wells to determine each well's actual gas and water production. If condensate is trucked out of the group separation and measurement point without further processing to a sales point, condensate production must be reported at the wellhead based on the condensate-gas ratio (CGR) from the well test. If liquid condensate is trucked to a gas plant for further processing, the condensate must be reported as a gas equivalent.

Multiwell Proration Battery

- Production from the wells in the battery is commingled prior to separation and measurement at a central location.

- Production from each well is not continuously measured, but is estimated based on periodic tests conducted at each well.

- The combined production of all wells in the proration battery must be separated and measured as single-phase components, and the resulting total actual battery gas volume (including GEV of condensate) and total actual battery water volume must be prorated back to the wells to determine each well's actual gas and water production.

- Gas wells that meet specific criteria applicable to Southeastern (SE) Alberta Shallow Gas Wells (see 7.2) may be grouped in a proration battery without specific approval. For those batteries, water volumes do not need to be prorated back to individual wells.

- Gas wells that do not meet the specific criteria applicable to SE Alberta Shallow Gas Wells may only be grouped in a proration battery if the conditions specified in Section 5: Site-specific Deviation from Base Requirements are met or if ERCB site-specific approval of an application is obtained.

4.2.3 Gas Gathering System

A facility consisting of pipelines used to move gas production from oil batteries, gas batteries, and/or other facilities to another facility (usually a gas plant) is considered to be a gas gathering system. It may include compressors, line heaters, dehydrators, and other equipment.

Inlet measurement usually consists of the battery or facility group measurement point.

Outlet measurement usually consists of the gas plant inlet measurement.

4.2.4 Gas Plant

A gas plant is a system or arrangement of equipment used for the extraction of H_2S, helium, ethane, NGLs, or other substances from raw gas. It does not include a wellhead separator, treater, dehydrator, or production facility that recovers less than 2 m^3/d of hydrocarbon liquids without using a liquid extraction process (e.g., refrigeration, desiccant). In addition, it does not include an arrangement of equipment that removes small amounts of sulphur (less than 0.1 tonne/day) through the use of nonregenerative scavenging chemicals that generate no H_2S or sulphur dioxide (SO_2).

Each plant inlet stream must have inlet separation and continuous measurement for all liquids and gas before commingling with other streams and must be used to report volume on the PRA for the plant receipt from upstream facilities (except for gas group batteries) and for plant balance. However, there are situations where the raw gas has been stripped of its liquid (not recombined downstream) and measured upstream of the plant site. If all streams entering a gas plant on the same gas gathering system are "dry" (the absence of free liquids), the gas plant inlet measurement may consist of the gas gathering system outlet measurement or battery group measurement.

Measurement of all gas deliveries out of the gas plant, such as sales, lease fuel for other facilities, flare and vent gas, acid gas disposition, and any volumes used internally, is required unless otherwise exempted by the ERCB. Monthly liquid inventory change must be accounted for and reported to the PRA. (See Figure 4.1.)
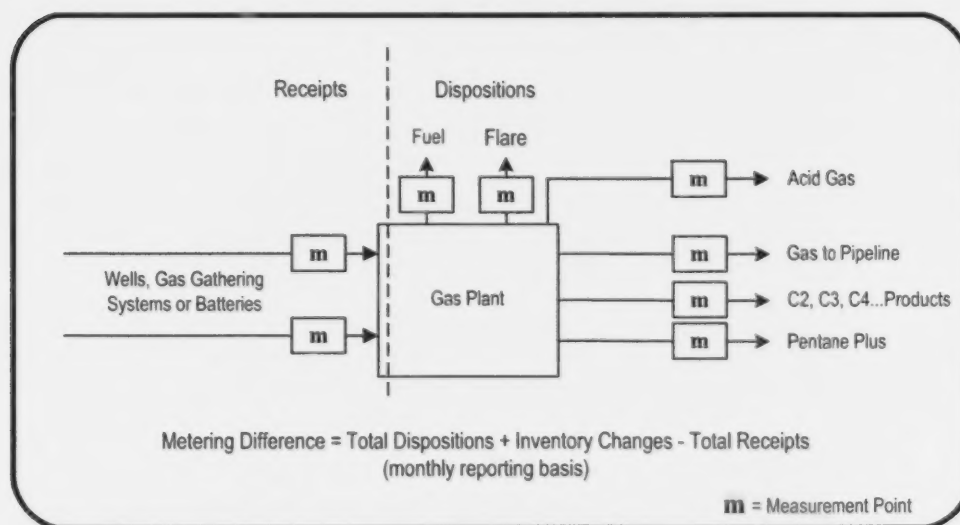
Figure 4.1. Typical gas plant measurement and reporting points

4.2.5   Gas Fractionation Plant

Condensate delivered to a gas fractionation plant licensed by the ERCB as a facility must be measured and reported in cubic metres by the operator of the fractionation facility as condensate received and reported, in accordance with existing ERCB requirements for trucked production.

4.3   Base Requirements for Gas Measurement

4.3.1   System Design and Installation of Measurement Devices

The design and installation of measurement systems must be in accordance with the following.

1) If an orifice meter system is used to measure gas, it must be designed and installed according to the applicable AGA Report #3: *Orifice Metering of Natural Gas and Other Related Hydrocarbon Fluids* (AGA3) listed in Table 4.1 below.

Table 4.1. Orifice meters design requirement (see detailed explanation in this section)

Meter run date of manufacture	Applicable AGA3 (API MPMS 14.3, Part 2) version
Before February 2003	AGA3 1991 or earlier meter run with upstream and/or downstream ID marking – may be reused or relocated for its designed application except to replace a meter where AGA3 2000 spec is required.
	Non-AGA meter run or run not marked with upstream or downstream ID – grandfathered for the existing volumetric throughput application; if relocated, it must be refurbished to AGA3 (1985) or later specification but cannot be used for sales/delivery point measurement.
After January 2003 (except for sales/delivery point meters measuring sales specification processed gas)	February 1991 or April 2000
All sales/delivery point meters measuring sales specification processed gas manufactured after January 2003	April 2000

When a meter such as a gas plant outlet meter is used to check sales/delivery point (royalty trigger point) measurement and is not normally used to report volumes to the ERCB, it does not require AGA3 April 2000 specification. However, when another gas source ties in to the sales pipeline between the check meter and the sales/delivery point (royalty trigger point) meter, the check meter could be used to report volumes to the ERCB. In this case, the AGA3 April 2000 specification is required if the meter is manufactured after January 2003, as shown in Figures 4.2 and 4.3, and the operator cannot replace the existing meter run with another meter run manufactured before February 2003 unless it meets the AGA3 April 2000 specification.
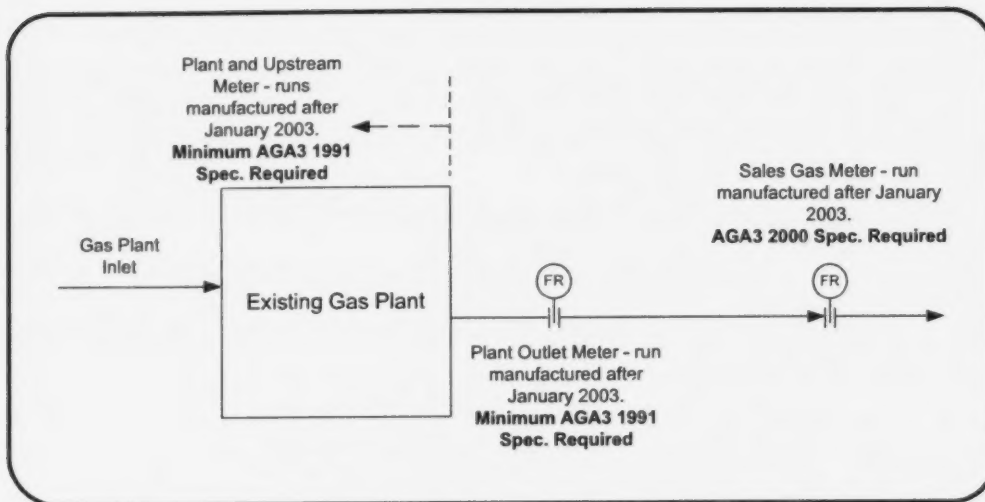
Figure 4.2
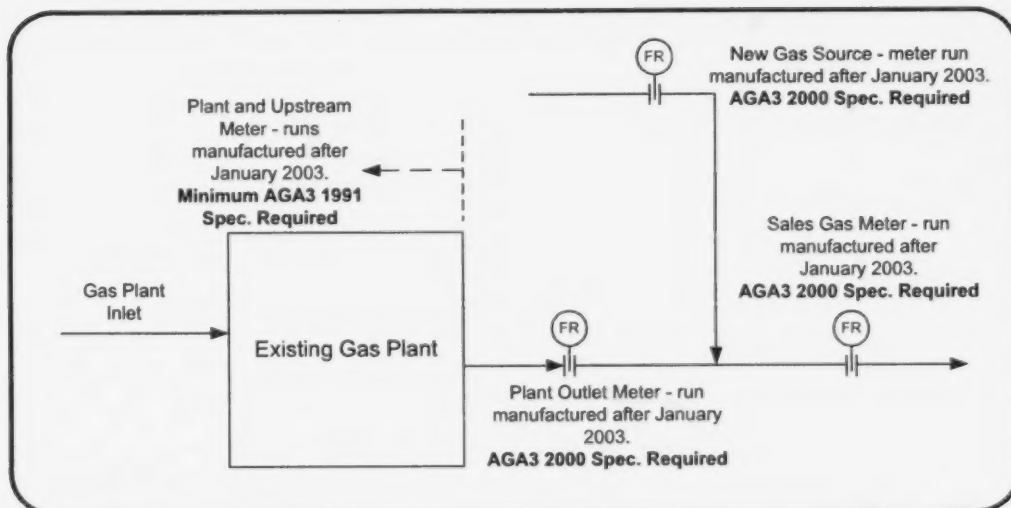
Figure 4.3

A permanently marked plate with the following information must be attached to each meter run and maintained in readable condition (not painted over or covered with insulation, etc.) for inspection:

- manufacturer's name
- serial number
- date of manufacture
- average upstream inside diameter (ID) of the meter run at 25.4 mm upstream of the orifice plate in millimetres to one decimal place (or to three decimal places if indicated in inches)

- AGA3 version/year (for new runs only after January 31, 2005) and/or AGA3 configuration for runs manufactured after January 2003 and not using the April 2000 specification

Meter runs that are manufactured before February 2003 and designed to the AGA3 1991 or earlier specifications complete with the upstream and/or downstream ID markings may be relocated or reused for the application they are designed for.

For existing in-service meter runs that are manufactured before February 2003 and are not designed to the AGA3 2000 or earlier specifications at the time of manufacture or not marked with upstream or downstream ID, nominal pipe ID can be used for flow calculations. These meter runs are grandfathered for the existing volumetric throughput. If new gas volumes are added to such an existing meter run or if a meter run is to be relocated, it must be inspected or refurbished to ensure that it meets the minimum of AGA3 1985 specifications, but it must not be used for sales/delivery point (royalty trigger point) measurement.

The orifice plate must be permanently marked with the plate bore in millimetres to two decimal places (or to three decimal places if indicated in inches), preferably within **6 mm** of the outside edge of the plate, to avoid interfering with normal flow if the marking creates a dent or protrusion on the plate surface.

The chart drive for a circular chart recorder used to measure gas well gas production or group oil battery gas production must not be more than 8 days per cycle unless the exception criteria specified in Section 5: Site-specific Deviation from Base Requirements are met or ERCB site-specific approval is obtained. A 24-hour chart drive is required for test gas measurement associated with single oil wells or Class 1 and 2 proration oil wells. An 8-day chart drive may be used for test gas measurement associated with Class 3 and 4 proration oil wells. If the mode of operation causes painting on the chart because of cycling or on/off flows, a 24-hour chart is required for any gas measurement point or EFM must be used.

Temperature measurement must be installed according to AGA3 specifications and the temperature must be determined in accordance with item 10 below.

2) Linear metering systems, such as a turbine or vortex meter, used to measure gas must be designed and installed according to the provisions of the 1985 or later editions of the AGA Report #7: *Measurement of Gas by Turbine Meters* (AGA7) or the manufacturer's recommendation.

   Temperature measurement is to be installed according to AGA7 (i.e., between one and five pipe diameter downstream of the meter) or the meter manufacturer's recommendation and the temperature must be determined as per item 8 below.

   The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections either on site (e.g., electronic correctors, electronic flow measurement) or at a later date (e.g., pressure and temperature charts).

3) Rotary metering systems must be designed and installed according to the provisions of the 1992 or later edition of the ANSI B109.3: *Rotary Type Gas Displacement Meters* or the manufacturer's recommendation.

   Install pressure taps not more than 20 pipe diameters upstream and downstream of the meter, to allow for measuring pressure drop across the meter and determining if the meter is over-ranging, if required. It is acceptable for the tap openings to be present within the

meter body. The upstream tap must be used for pressure measurement and must be reading the metering pressure (i.e., there must be no pressure restriction between the tap and the meter, such as a regulator).

The meter must be equipped with a non-reset counter. This can be mechanical or electronic.

Install temperature measurement according to the meter manufacturer's recommendation or less than 20 pipe diameters downstream of the meter, with no restrictions between the meter and the temperature probe. The temperature must be determined as per item 8 below.

The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections either on site (e.g., electronic correctors, electronic flow measurement) or at a later date (e.g., pressure and temperature charts).

4) Diaphragm displacement metering systems must be designed and installed according to the provisions of the 1992 or later edition of the ANSI B109.1: *Diaphragm Type Gas Displacement Meters (up to 500 cubic feet/hour capacity)*, or ANSI B109.2: *Diaphragm Type Gas Displacement Meters (over 500 cubic feet/hour capacity)*, and/or the manufacturer's recommendation.

   Other conditions are the same as for the rotary meter above.

5) Venturi or flow nozzle type of metering systems must be installed according to the provisions of the 1991 or later edition of the ISO Standard 5167: *Measurement of fluid flow by means of orifice plates, nozzles and venturi tubes inserted in circular cross-section conduits running full* (ISO 5167) or the meter manufacturer's recommendation. The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections either on site or at a later date.

6) Ultrasonic metering systems must be designed and installed according to the provisions of the 1998 or later editions of *AGA Report No. 9: Measurement of Gas by Multipath Ultrasonic Meters* (AGA9). The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections.

7) Coriolis mass metering systems must be designed and installed according to the provisions of the latest edition of *AGA Report No. 11: Measurement of Natural Gas by Coriolis Meter*. The installation must include instrumentation that allows for continuous temperature correction.

8) If metering systems other than those listed above, such as v-cones or wedge meters, are used to measure gas, they must be installed according to applicable industry accepted standards or the meter manufacturer's recommendation. The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections (where required) either on site or at a later date.

9) Any electronic gas measurement system must be designed and installed according to the requirements as stated in Section 4.3.3.2. Any EFM system designed and installed in accordance with API MPMS, Chapter 21.1 is considered to have met the audit trail and reporting requirements. However, the performance evaluation is still required in accordance with Section 4.3.3.2. All EFM devices must have a continuous temperature reading for flow calculation.

10) The flowing gas temperature must be measured and recorded according to Table 4.2.

11) Any measurement under vacuum conditions must have absolute pressure measurement to accurately measure the static pressure.

Table 4.2. Temperature reading frequency table for gas measurement

Minimum temperature reading frequency	Criteria or events
Continuous	Sales/delivery points (royalty trigger point) and/or EFM devices
Daily	$> 16.9 \ 10^3 \ \mathrm{m}^3/\mathrm{d}$
Weekly	$\leq 16.9 \ 10^3 \ \mathrm{m}^3/\mathrm{d}$
Daily	a) Production (proration) volume testing, or b) Nonroutine or emergency flaring and venting

Note that the temperature-measuring element must be installed near the meter such that it will be sensing the flowing gas stream temperature. That is, the licensee cannot use the surface temperature of the piping or use a thermowell location where there is normally no flow. A meter equipped with a temperature compensation device is considered to have continuous temperature measurement.

4.3.2 Volumetric Calculations

The gas volume calculations comply if the following requirements are met.

- If an orifice meter is used to measure gas, the licensee must use the 1985 or later editions of the AGA3 to calculate the gas volumes.

- If a positive displacement meter or a linear type of meter, such as a turbine, ultrasonic, or vortex meter, is used to measure gas, volumes must be calculated according to the provisions of the 1985 or later editions of the AGA7. Corrections for static pressure, temperature, and compressibility are required.

- If a venturi or flow nozzle type of meter is used to measure gas, volumes must be calculated according to the provisions of the 1991 or later edition of the ISO 5167 or the meter manufacturer's recommended calculation procedures.

- If a Coriolis mass meter is used to measure gas, volumes must be calculated from the measured mass flow and the base density derived from a representative gas sample analysis, including corrections for compressibility. The flowing density measured by the Coriolis mass meter is of insufficient accuracy in a gas application and must not be used to compute volumes.

- If meter types other than those listed above, such as v-cones or wedge meters, are used to measure gas, volumes must be calculated according to the applicable industry accepted standard or the meter manufacturer's recommendation.

- If condensate production from a gas well is required to be reported as a gas equivalent volume, the calculation of the gas equivalent factor must be performed in accordance with the methodologies outlined in Section 8.3. The following are the general requirements:

    - The gas equivalent volume (GEV) is to be determined based on the latest condensate sample analysis.

- The gas equivalent volume can be determined using the volume fractions, mole fractions, or mass fractions of the condensate analysis.

- The gas equivalent volume can be determined using all of the individual components in the condensate analysis, *or* the C_5 and/or heavier components in the sample can be grouped as C_{5+}, C_{6+}, C_{7+} or other heavier component groups. If the heavier components are grouped, the gas equivalent factor for the grouped components must be calculated using the molecular weight and/or relative density of the grouped components.

- Correction for deviation from the Ideal Gas Laws for compressibility is to be based on equations published in the November 1992, second edition of the AGA Transmission Measurement Committee Report No. 8 (AGA8): *Compressibility Factors of Natural Gas and Other Related Hydrocarbon Gases* or one of the methods listed in the section below. For EFM systems installed before 1994 with software or hardware limitations incompatible with the second edition of AGA8, an earlier version can be used.

Compressibility Factors Used in Gas Volume Calculations

Produced or injected gas volume measurements must be corrected for pressure, temperature, gas composition, and the compressibility of the natural gas.

The AGA8[1] (1992) or Redlich-Kwong with Wichert-Aziz sour gas corrections method should be used for the calculation of the compressibility factors. However, other methods can also be used, provided that the licensee documents the reason for their use. Other methods that could be used are

- Pitzer et al. with Wichert-Aziz sour gas corrections

- Dranchuk, Purvis, Robinson with Wichert-Aziz sour gas corrections (Standing and Katz)

- Dranchuk, Abou-Kassam with Wichert-Aziz sour gas corrections (Starling)

- Hall, Yarborough with Wichert-Aziz sour gas corrections

The ERCB will also accept the use of methods other than those mentioned above. If others are used, a suitable reference and comparison to the AGA8 (1992) method or to experimental results and the justification for use must be documented and provided to the ERCB for inspection on request.

The AGA8 publication includes several approaches for estimating the properties of natural gas for use in the AGA8 calculation. The full compositional analysis (Detail) method must be used rather than the less accurate partial composition method.

If paper charts are used, the compressibility factor should be calculated at least once for each gas chart cycle. Flow computers and other EFM systems used for gas measurement must calculate and update the compressibility (or supercompressibility) factor at a minimum of once every five minutes, whenever the gas composition is updated, or whenever the average pressure or temperature changes by more than ± 0.5% from the previous average 5-minute value used for calculation.

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[1] See Section 4.4: References for complete bibliographical details for these citations.

Physical Properties of Natural Gas Components

The ERCB adopts the physical properties contained in the most recent edition of the Gas Processors Suppliers Association (GPSA) *SI Engineering Data Book*[1] or the Gas Processors Association (GPA) *2145*[1] publication, whichever is the most current. The licensee must ensure that it is using the up-to-date list and, if necessary, update its data. If an EFM system does not have the capability to accept updated physical constants, then the existing set of physical constants may be used; however, that type of EFM system must not be used for measurement of processed gas that meets sales specifications. For standards, such as AGA8, that have imbedded physical constants different in value from those in GPA 2145 or GPSA *SI Engineering Data Book*, changes to such standards are not required unless they are made by the standard association.

4.3.3 Production Data Verification and Audit Trail

4.3.3.1 General

The field data, records, and any calculations or estimations including EFM relating to ERCB-required production data submitted to the PRA must be kept for inspection on request. The reported data verification and audit trails must be in accordance with the following:

1) When a bypass around a meter is opened or when, for any reason, gas does not reach the meter or the recording device, a reasonable estimate of the unmetered volume must be determined, the method used to determine the estimate must be documented, and a record of the event must be made.

2) A record must be maintained that identifies the gas stream being metered, the measurement devices, and all measurements, inputs, times, and events related to the determination of gas volumes. (See Section 4.3.4: Operations for more detail on orifice chart recorders.) If EFM is used, the required data must be collected and retained according to Section 4.3.3.2.

3) Test records: Any documentation produced in the testing or operation of metering equipment that affects measured volumes. This includes the record containing volume verification and calibration measurements for all secondary and tertiary devices.

4) When a gas metering error is discovered, the licensee of the facility must immediately correct the cause of the error and submit amended monthly production reports to the PRA to correct all affected gas volumes.

5) All flared and vented gas must be reported as described in ERCB *Directive 007: Production Accounting Handbook*. Incinerated gas must be reported as "flared" gas if an incinerator is used in place of a flare stack. This would not apply to acid gas streams at a gas plant that are incinerated as part of normal operations; in such cases, the incinerated acid gas would be reported as shrinkage, not as flared.

6) Whenever possible, the licensee must report gas as fuel, flared, or vented as occurring at the location where the fuel use, flaring, or venting took place. This will allow industry and ERCB staff to match flaring or venting that is observed in the field with that reported. When the fuel usage, flaring, or venting location is within a gas gathering system but is not at a licensed entity,

   a) it should be reported as an activity associated with the closest licensed facility (e.g., compressor) within the gas gathering system, or

b) if there is no applicable licensed facility within the gas gathering system, it should be reported as an activity associated with the gas gathering system itself.

7) Licensees must not prorate or allocate flared and vented volumes that occur at a facility to other upstream facilities and/or well locations.

8) Dilution gas, purge gas, or gas used to maintain a minimum heating value of the flared or incinerated gas is to be reported as fuel. The reported total flare volume must exclude any of these fuel volumes.

9) Production hours for gas wells designed to operate on an on/off cycle basis, such as intermittent timers, pump-off controls, plunger lifts, manual on/off, or pumpjacks, that are "operating normally and as designed on repeated cycles" and where part of the operation involves shutdown of pump equipment and/or shut-in as part of the repeated cycles are to be considered on production even when the wells are not flowing. Physical well shut-ins (not as part of a repeated cycle) and emergency shutdowns (ESDs) are considered down time. The operation personnel have to make a decision based on the operating environment in other situations where the wells are not shut in but may or may not have production.

10) All gas usage, such as for instrumentation, pumps, and heating, must be reported as fuel use on a per-site basis, even if it is vented afterwards. The volume must be measured on a per site basis if over $0.5 \ 10^3 \ \text{m}^3/\text{d}$ or may be estimated if not over $0.5 \ 10^3 \ \text{m}^3/\text{d}$ (see Figure 1.11). If there are multiple reporting facilities on the same site, the fuel use has to be separately reported to each individual facility but there must be overall site fuel measurement.

4.3.3.2 Electronic Flow Measurement (EFM) for Gas

An EFM system is defined as any flow measurement and related system that collects data and performs flow calculations electronically. If it is part of a Distributed Control System (DCS), Supervisory Control and Data Acquisition system (SCADA) or Programmable Logic Controller system (PLC), only the EFM portion has to meet the requirements in this section. All EFM systems approved under *Guide 34* may continue as approved.

The following systems are not defined as an EFM:

- any meter with an electronic totalizer or pulse counter that does not perform flow calculations (with or without built-in temperature compensation)

- a Remote Terminal Unit (RTU) that transmits any data other than flow data and does not calculate flow

4.3.3.2(a) Base Requirements for EFM

If an EFM is used to calculate volumes for ERCB accounting purpose, the licensee must be able to verify that it is performing within the ERCB target limits defined in this section.

When any parameter that affects the flow calculation is changed, such as orifice plate size, meter factor, fluid analysis, or transmitter range, a signoff procedure or an event log must be set up to ensure that the change is made in the EFM system. All data and reports must be retained for a minimum of 12 months.

Hardware and software requirements:

- The memory on board the EFM must allow for at least 32 days of storage of the required flow data before being overwritten or erased.

- The EFM must be equipped with its own on-board battery to protect the memory in the event of a power failure.

- The system must have various levels of system security, with the highest level of access to the program restricted to authorized people.

- The communication system must use a data integrity error-checking routine to ensure that the data transmitted are correct.

- The super-compressibility must be calculated and updated every five minutes or every flow calculation point, whichever occurs first.

- The EFM must be set to alarm on high and low differential pressure, over-range of any end devices, low power, and communication failures.

- Any changes made to the data or any manually entered values that affect the flow calculation must be flagged so it is clear that these are estimated, not actual, readings.

- This flagging must carry through to values calculated from the data,

- Slower sampling or integration frequencies than stated in *API-MPMS*, Chapter 21, Clauses 1.4.2.1, 1.4.2.2, and 1.4.3.1, may be used if it can be demonstrated using the Rans Methodology that uncertainties are still being met,

- Hourly quantity transaction records as specified in *API-MPMS*, Chapter 21, Clauses 1.6.2.2 and 1.6.2.4, are not required. Only daily transaction records are required.

4.3.3.2(b)   Performance Evaluation

A performance evaluation test must be completed within **two weeks** after the EFM is put into service and **immediately** after any change to the computer program or algorithms that affects the flow calculation; it must be documented for ERCB audit on request. For existing EFM systems, the ERCB encourages licensees to conduct their own performance evaluations. A performance evaluation must be conducted and submitted for ERCB audit on request. The ERCB considers either one of the following methods acceptable for performance evaluation:

1) Conduct a performance evaluation test on the system by inputting known values of flow parameters into the EFM to verify the volume calculation, coefficient factors, and other parameters. The first **seven** test cases included in this section are for gas orifice meters (AGA3 flow calculations), each with different flow conditions and gas properties. **Test Case 8** is for the AGA7 flow calculation for positive displacement or linear meters. Other manufacturers' recommended equations can also be used to evaluate the EFM performance. The seven AGA3 test cases could also be used to evaluate any compressibility or supercompressibility factors used in other flow calculations using the same gas composition, pressure, and temperature in the calculation as inputs.

2) Evaluate the EFM calculation accuracy with a flow calculation checking program that performs within the target limits for all the factors and parameters listed in the test cases below. A snapshot of the instantaneous flow parameters and factors, flow rates, and configuration information is to be taken from the EFM and input into the checking program. If the instantaneous EFM flow parameters, factors, and flow rates are not updated simultaneously, multiple snapshots may have to be taken to provide a representative evaluation.

Note that some DCS or other control systems have built-in and/or manual input of pressure and temperature for flow calculations. Since the pressure and temperature are not continuously updated, they are not acceptable for ERCB accounting and reporting purposes unless ERCB approval is obtained.

The volumetric flow rate (Q) obtained from a performance evaluation test must agree to within ±0.25% of those recorded on the sample test cases or other flow calculation checking programs. If the ± 0.25% limit is exceeded, the EFM must be subjected to a detailed review of the calculation algorithm to resolve the deviation problem. For gas orifice meters, if no AGA3 factor or parameter outputs are available, the acceptable volumetric gas flow rate limit is lowered to ± 0.15%.

Test Cases 1 to 7 for Verification of Orifice Meter Gas Flow Calculation Programs

The ERCB has developed test cases to verify that the EFM system correctly calculates gas flow rates from orifice meters. The seven test cases were calculated on the following basis:

- They are for flange taps only.

- The atmospheric pressure is assumed to be 93.08 kPa(a) (13.5 psia).

- The heaviest carbon component was assumed to be normal heptane.

- The ideal gas relative density was converted to the real gas relative density.

- The same static pressure value is used for pressure taps that are located upstream (U/S) or downstream (D/S) of the orifice plate.

- The AGA3 (1985) results were calculated based on upstream conditions (for both upstream and downstream static pressure tap) in imperial units (the Y_2 factor is also provided for reference). The metric conversion factor for the calculated gas volume is 0.02831685. The compressibility factors were calculated using the Redlich-Kwong (RK) equation with the Wichert-Aziz correction for sour gas.

- The AGA3 (1990) results were calculated using the Detail AGA8 (1992) compressibility factor calculation and using the upstream expansion factor Y_1, as recommended by the AGA3 (1990), Part 1, Section 1.8, even though the pressure tap may be downstream of the orifice plate. (The Y_2 factor is also provided for reference when applicable.)

- The orifice plate material is assumed to be 316 stainless steel and the meter run to be carbon steel at reference temperature of 20°C, isentropic exponent (k) = 1.3, viscosity = 0.010268 centipoise.

- The base conditions (101.325 kPa[abs] and 15°C) are used in the calculated temperature base factor (F_{tb}) and pressure base factor (F_{pb})

Test Case 8 for Verification of AGA7 Gas Flow Calculation Programs

The ERCB has developed a test case to verify that the EFM system correctly calculates gas flow rates using the AGA7 equations. The test case was calculated on the following basis:

- The heaviest carbon component was assumed to be normal heptane.

- The compressibility factors were calculated using the Detail AGA8 (1992) or the Redlich-Kwong (RK) equation with the Wichert-Aziz correction for sour gas.

Table 4.3. Allowable deviation limits for the AGA3 (1985) equation

AGA3 (1985) factors	Allowed deviation limit from test cases
Y, F_a, F_r, and F_{tf}	± 0.01%
F_b	± 0.1%
F_{gr}, F_{pv}	± 0.2%
Q	± 0.25% or ± 0.15% without the above factors

Table 4.4. Allowable deviation limits for the AGA3 (1990) equation

AGA3 (1990) factors	Allowed deviation limit from test cases
Y_1, and E_v	± 0.01%
C_d and Z_b	± 0.1%
Z_f	± 0.2%
Q	± 0.25% or ± 0.15% without the above factors

Table 4.5. Allowable deviation limits for the AGA7 equation

AGA7 factors	Allowed deviation limit from test cases
F_{pm} (flowing pressure) and F_{tm} (flowing temperature)	± 0.1%
S (compressibility)	± 0.2%
Q	± 0.25% or ± 0.15% without the above factors

TEST CASE 1 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0184	iC_4	-	0.0081
CO_2	-	0.0000	nC_4	-	0.0190
H_2S	-	0.0260	iC_5	-	0.0038
C_1	-	0.7068	nC_5	-	0.0043
C_2	-	0.1414	C_6	-	0.0026
C_3	-	0.0674	C_7	-	0.0022

Ideal gas relative density - 0.7792

Meter Data (flange taps)

Meter run	I.D.	-	52.370 mm (2.0618 inches)
Orifice	I.D.	-	9.525 mm (0.375 inches)

Flow Data (24 hr)

Static pressure	-	2818.09 kPa(a) (408.73 psia)
Differential pressure	-	10.2000 kPa (40.9897 inches H_2O)
Flowing temperature	-	57.0°C (134.600°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap
F_b	28.4286	28.4286
Y_1	0.9989	0.9989
Y_2	N/A	1.0007
F_{tb}	0.9981	0.9981
F_{gr}	1.1308	1.1308
F_a	1.0012	1.0012
F_r	1.0006	1.0006
F_{pb}	1.0023	1.0023
F_{tf}	0.9351	0.9351
F_{pv}	1.0360	1.0361
C'	31.175	31.179
Q	2.7422	2.7475 10^3 m^3/24 hr

AGA3 (1990)

Factors	U/S Tap	D/S Tap
C_d	0.5990	0.5990
Y_1	0.9989	0.9989
Y_2	N/A	1.0007
E_v	1.0005	1.0005
Z_b	0.9959	0.9959
Z_f	0.9280	0.9277
Q	2.7478	2.7531 10^3 m^3/24 hr

TEST CASE 2 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0156	iC_4	-	0.0044
CO_2	-	0.0216	nC_4	-	0.0075
H_2S	-	0.1166	iC_5	-	0.0028
C_1	-	0.7334	nC_5	-	0.0024
C_2	-	0.0697	C_6	-	0.0017
C_3	-	0.0228	C_7	-	0.0015

Ideal gas relative density - 0.7456

Meter Data (flange taps)

Meter run	I.D.	-	102.26 mm (4.026 inches)
Orifice	I.D.	-	47.625 mm (1.875 inches)

Flow Data (24 hr)

Static pressure	-	9100.94 kPa(a) (1319.98 psia)
Differential pressure	-	11.0000 kPa (44.2046 inches H_2O)
Flowing temperature	-	50.0°C (122.0°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap
F_b	733.697	733.697
Y_1	0.9996	0.9996
Y_2	N/A	1.0002
F_{tb}	0.9981	0.9981
F_{gr}	1.1564	1.1564
F_a	1.0010	1.0010
F_r	1.0002	1.0002
F_{pb}	1.0023	1.0023
F_{tf}	0.9452	0.9452
F_{pv}	1.1072	1.1073
C'	888.905	889.000
Q	145.93	146.03 10^3 m^3/24 hr

AGA3 (1990)

Factors	U/S Tap	D/S Tap
C_d	0.6019	0.6019
Y_1	0.9996	0.9996
Y_2	N/A	1.0003
E_v	1.0244	1.0244
Z_b	0.9967	0.9967
Z_f	0.8098	0.8097
Q	146.08	146.18 10^3 m^3/24 hr

TEST CASE 3 (for AGA3 Flow Calculations)

Gas Analysis

N_2	–	0.0500		iC_4	–	0.0000
CO_2	–	0.1000		nC_4	–	0.0000
H_2S	–	0.2000		iC_5	–	0.0000
C_1	–	0.6000		nC_5	–	0.0000
C_2	–	0.0500		C_6	–	0.0000
C_3	–	0.0000		C_7	–	0.0000

Ideal gas relative density - 0.8199

Meter Data (flange taps)

Meter run	I.D.	–	590.55 mm (23.250 inches)
Orifice	I.D.	–	304.80 mm (12.000 inches)

Flow Data (24 hr)

Static pressure	–	10342.14 kPa(a) (1500.00 psia)
Differential pressure	–	22.1600 kPa (89.0522 inches H_2O)
Flowing temperature	–	60.0°C (140.0°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap
F_b	30429.66	30429.66
Y_1	0.9993	0.9993
Y_2	N/A	1.0004
F_{tb}	0.9981	0.9981
F_{gr}	1.1028	1.1028
F_a	1.0013	1.0013
F_r	1.0001	1.0001
F_{pb}	1.0023	1.0023
F_{tf}	0.9309	0.9309
F_{pv}	1.1076	1.1078
C'	34636.6	34643.21
Q	8603.19	8614.04 10^3 m³/24 hr

AGA3 (1990)

Factors	U/S Tap	D/S Tap
C_d	0.6029	0.6029
Y_1	0.9993	0.9993
Y_2	N/A	1.0004
E_v	1.0375	1.0375
Z_b	0.9968	0.9968
Z_f	0.8216	0.8213
Q	8564.77	8575.48 10^3 m³/24 hr

TEST CASE 4 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0029	iC_4	-	0.0000
CO_2	-	0.0258	nC_4	-	0.0000
H_2S	-	0.0000	iC_5	-	0.0000
C_1	-	0.9709	nC_5	-	0.0000
C_2	-	0.0003	C_6	-	0.0000
C_3	-	0.0001	C_7	-	0.0000

Ideal gas relative density - 0.5803

Meter Data (flange taps)

Meter run	I.D.	-	146.36 mm (5.7622 inches)
Orifice	I.D.	-	88.900 mm (3.500 inches)

Flow Data (24 hr)

Static pressure	-	9839.99 kPa(a) (1427.17 psia)
Differential pressure	-	6.6130 kPa (26.575 inches H_2O)
Flowing temperature	-	22.35°C (72.23°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap
F_b	2694.965	2694.97
Y_1	0.9998	0.9998
Y_2	N/A	1.0001
F_{tb}	0.9981	0.9981
F_{gr}	1.3116	1.3116
F_a	1.0001	1.0001
F_r	1.0002	1.0002
F_{pb}	1.0023	1.0023
F_{tf}	0.9884	0.9884
F_{pv}	1.0843	1.0843
C'	3790.16	3790.31
Q	501.64	501.82 10^3 m³/24 hr

AGA3 (1990)

Factors	U/S Tap	D/S Tap
C_d	0.6047	0.6047
Y_1	0.9998	0.9998
Y_2	N/A	1.0001
E_v	1.0759	1.0759
Z_b	0.9980	0.9980
Z_f	0.8425	0.8425
Q	503.44	503.63 10^3 m³/24 hr

TEST CASE 5 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0235	iC_4	-	0.0088
CO_2	-	0.0082	nC_4	-	0.0169
H_2S	-	0.0021	iC_5	-	0.0035
C_1	-	0.7358	nC_5	-	0.0031
C_2	-	0.1296	C_6	-	0.0014
C_3	-	0.0664	C_7	-	0.0007

Ideal gas relative density - 0.7555

Meter Data (flange taps)

Meter run	I.D.	-	154.05 mm (6.0650 inches)
Orifice	I.D.	-	95.250 mm (3.750 inches)

Flow Data (24 hr)

Static pressure	-	2499.9 kPa(a) (362.58 psia)
Differential pressure	-	75.000 kPa (301.395 inches H_2O)
Flowing temperature	-	34.0°C (93.2°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap
F_b	3111.24	3111.24
Y_1	0.9894	0.9897
Y_2	N/A	1.0044
F_{tb}	0.9981	0.9981
F_{gr}	1.1485	1.1485
F_a	1.0005	1.0005
F_r	1.0001	1.0001
F_{pb}	1.0023	1.0023
F_{tf}	0.9695	0.9695
F_{pv}	1.0382	1.0394
C'	3561.90	3567.34
Q	800.22	813.37 $10^3 \, m^3/24 \, hr$

AGA3 (1990)

Factors	U/S Tap	D/S Tap
C_d	0.6042	0.6041
Y_1	0.9894	0.9897
Y_2	N/A	1.0044
E_v	1.0822	1.0822
Z_b	0.9962	0.9962
Z_f	0.9240	0.9217
Q	799.83	813.00 $10^3 \, m^3/24 \, hr$

TEST CASE 6 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0268	iC_4	-	0.0123
CO_2	-	0.0030	nC_4	-	0.0274
H_2S	-	0.0000	iC_5	-	0.0000
C_1	-	0.6668	nC_5	-	0.0000
C_2	-	0.1434	C_6	-	0.0180
C_3	-	0.1023	C_7	-	0.0000

Ideal gas relative density - 0.8377

Meter Data (flange taps)

Meter run	I.D.	-	52.500 mm (2.0669 inches)
Orifice	I.D.	-	19.050 mm (0.750 inches)

Flow Data (24 hr)

Static pressure	-	2506.33 kPa(a) (363.50 psia)
Differential pressure	-	17.0500 kPa (68.5171 inches H_2O)
Flowing temperature	-	7.2°C (44.96°F)

Gas Volume Result

AGA3 (1985)				**AGA3 (1990)**		
Factors	**U/S Tap**	**D/S Tap**		**Factors**	**U/S Tap**	**D/S Tap**
F_b	115.138	115.138		C_d	0.6005	0.6005
Y_1	0.9978	0.9978		Y_1	0.9978	0.9978
Y_2	N/A	1.0012		Y_2	N/A	1.0012
F_{tb}	0.9981	0.9981		E_v	1.0088	1.0088
F_{gr}	1.0902	1.0902		Z_b	0.9951	0.9951
F_a	0.9996	0.9996		Z_f	0.8588	0.8578
F_r	1.0003	1.0003		Q	14.687	14.746 10^3 m^3/24 hr
F_{pb}	1.0023	1.0023				
F_{tf}	1.0148	1.0148				
F_{pv}	1.0708	1.0714				
C'	136.15	136.22				
Q	14.602	14.660 10^3 m^3/24 hr				

TEST CASE 7 (for AGA3 Flow Calculations)

Gas Analysis

N_2	-	0.0070	iC_4	-	0.0062
CO_2	-	0.0400	nC	-	0.0090
H_2S	-	0.0000	iC_5	-	0.0052
C_1	-	0.8720	nC_5	-	0.0016
C_2	-	0.0340	C_6	-	0.0000
C_3	-	0.0250	C_7	-	0.0000

Ideal gas relative density - 0.6714

Meter Data (flange taps)

Meter run	I.D.	-	52.500 mm (2.0669 inches)
Orifice	I.D.	-	12.70 mm (0.50 inches)

Flow Data (24 hr)

Static pressure	-	299.92 kPa(a) (43.50 psia)
Differential pressure	-	6.3455 kPa (25.5 inches H_2O)
Flowing temperature	-	1.67°C (35°F)

Gas Volume Result

AGA3 (1985)

Factors	U/S Tap	D/S Tap	
F_b	50.523	50.523	
Y_1	0.9933	0.9935	
Y_2	N/A	1.0039	
F_{tb}	0.9981	0.9981	
F_{gr}	1.2190	1.2190	
F_a	0.9994	0.9994	
F_r	1.0018	1.0018	
F_{pb}	1.0023	1.0023	
F_{tf}	1.0250	1.0250	
F_{pv}	1.0035	1.0036	
C'	63.013	63.029	
Q	1.4263	1.4416	$10^3\,m^3$/24 hr

AGA3 (1990)

Factors	U/S Tap	D/S Tap	
C_d	0.6006	0.6006	
Y_1	0.9933	0.9934	
Y_2	N/A	1.0039	
E_v	1.0017	1.0017	
Z_b	0.9973	0.9973	
Z_f	0.9905	0.9903	
Q	1.4335	1.4489	$10^3\,m^3$/24 hr

TEST CASE 8 (for AGA7 Flow Calculations)

Gas Analysis

N_2	-	0.0268	iC_4	-	0.0123
CO_2	-	0.0030	nC_4	-	0.0274
H_2S	-	0.0000	iC_5	-	0.0000
C_1	-	0.6668	nC_5	-	0.0000
C_2	-	0.1434	C_6	-	0.0180
C_3	-	0.1023	C_7	-	0.0000

Flow Data (24 hr)

Uncorrected volume   - $128.0\ 10^3 m^3$
Static pressure   -   2506.33 kPa(a) (363.50 psia)
Flowing temperature   -   7.2°C (44.96°F)

Gas Volume Result

AGA7 (Volumetric Flow)
Factors

F_{pm} 24.6784

F_{pb} 1.0023

F_{tm} 1.0298

F_{tb} 0.9981

Using AGA8 compressibility equations,
S   1.1588
Q $3770.9\ 10^3\ m^3/24\ hr$

Using RK compressibility equations,
S   1.1467
Q $3731.6\ 10^3\ m^3/24\ hr$

EFM Reports

The required information on each report must be stored using electronic/magnetic (not necessarily on the EFM) or printed media and can exist individually on different formats or reports and generated on demand for audit, as follows:

- **Daily** for daily report required data
- **Monthly** for monthly report required data
- Event and alarm logs at regular intervals before information is overwritten
- Meter reports generated on request for audit

The Daily Report

The daily report must include

- meter identification
- daily accumulated flow, with indicating flags for estimated flows made by the system or by the operation personnel and alarms that have occurred for over-ranging of end devices
- hours on production or hours of flow (specify)
- flow data audit trail – include at least one of the following:
  - instantaneous values for flow rate, differential pressure (if applicable), static pressure, and temperature taken at the same time each day, or
  - average daily values for differential pressure (if applicable), static pressure, and temperature, or
  - hourly accumulated flow rate and average hourly values for differential pressure (if applicable), static pressure, and temperature

Existing EFM systems that predate *Guide 34* approvals and do not have any of the above audit trail capabilities, and cannot develop the capability due to system limitations, should be evaluated for upgrading, especially when new production is tied into the facilities. The ERCB may request upgrades, where audit/inspection results indicate they are warranted.

The Monthly Report

This report is for the entire system, providing data for each measurement point. It is to contain the following at each measurement point as applicable:

- monthly cumulative flow
- flags indicating any change made to flow volumes
- total hours on production or hours of flow (specify)

The Meter Report

The meter report details the configuration of each meter and flow calculation information. These values are used as part of the "audit trail" to confirm that the flow calculation is functioning correctly. Without them there is no way of verifying the accuracy of the system. The meter report must include the following as applicable and be produced on demand:

1) Instantaneous flow data
   - Instantaneous flow rate

- Instantaneous static pressure
- Instantaneous differential pressure
- Instantaneous flowing temperature
- Instantaneous relative density (if live)
- Instantaneous compressibility (if live)
- Instantaneous gas component (if live)
- Optional: instantaneous (AGA3) factors (see the orifice meter test cases above for output information)

2) Current configuration information for differential meters or other types of meters, whichever are applicable:

- meter identification
- date and time
- contract hour
- atmospheric pressure
- pressure base (unless fixed)
- temperature base (unless fixed)
- meter tube reference inside diameter
- orifice plate reference bore size
- static pressure tap location
- orifice plate material
- meter tube material
- calibrated static pressure range
- calibrated differential pressure range
- calibrated temperature range
- high/low differential cutoff
- relative density (if not live)
- compressibility (if not live)
- gas components (if not live)
- meter factor and/or k factor
- effluent correction factor

The Event Log

This log is used to note and record exceptions and changes to the flow parameter, configuration, programming, and database affecting flow calculations, such as, but not limited to,

- orifice size change
- transmitter range change

- date of gas/liquid analysis update
- algorithm changes
- meter factor, k factor, or effluent correction factor changes
- other manual inputs

The Alarm Log

The alarm log includes any alarms that may have an effect on the measurement accuracy of the system. The time of each alarm condition and the time of clearing of each alarm must be recorded. Alarms to be reported must include, but are not limited to,

- master terminal unit failure
- remote terminal unit failures
- communication failures
- low-power warning
- high differential pressure (for differential measurement devices)
- high/low volumetric flow rate (for other types of measurement)
- over-ranging of end devices

4.3.4   Operations—Base Requirements for Acceptable Gas Measurement and Reading of Gas Charts

4.3.4.1 General Operations

- Accounting meters using differential pressure sensing devices must be equipped with full port valves at the metering tap on the sensing lines. The valves must be the same size as the sensing lines (12.7 mm minimum for meter runs 102 mm in diameter or larger, and 9.5 mm minimum for meter runs less than 102 mm). All metering design and installation must ensure that the sensing line diameter does not change from the sensing tap valve to the manifold for delivery point, group point, and sales point measurement.

- Sharing of metering taps by multiple differential pressure devices is not allowed if it will cause increased measurement uncertainty, such as painting or spiking charts, or under pulsation conditions.

- A separate set of valve manifolds must be used for each device.

- The orifice meter must be in good operating condition.

- Chart recorders, sensing lines, and other piping must be in good operating condition and suitably winterized to prevent them from freezing and disrupting measurement.

- Sensing lines must be self-draining towards the sensing taps to prevent liquid from being trapped in the line and disrupting measurement accuracy from **February 2, 2009, onward** if they do not meet the exception criteria for changes in sensing line diameter below.

4.3.4.1(a) Exceptions

Sensing Line Tap Valves and Changes in Sensing Line Diameter

Grandfathering of existing differential pressure-sensing tap valves for installation before May 7, 2007, is granted without application **unless**

- the metering device is being upgraded, refurbished, and commissioned within a new application or relocated, or
- the metering device does not meet the single point uncertainty limit, as detailed in Section 1: Standards of Accuracy, or
- the metering point is subject to noticeable pulsation effects, such as physical vibration or audible flow noise, or is downstream of a reciprocal compressor on the same site, or
- the metering point is at a delivery point, group point, sales point (royalty trigger point), or custody transfer point.

Grandfathering of changes in sensing line diameter from the sensing tap to the manifold, such as drip pots, installed **before February 2, 2009,** is granted without application **unless**

- the metering device does not meet the single point uncertainty limit, as detailed in Section 1: Standards of Accuracy, or
- the metering point is subject to noticeable pulsation effects, such as physical vibration or audible flow noise, or is downstream of a reciprocal compressor on the same site, or
- the metering point is at a delivery point, group point, sales point (royalty trigger point), or custody transfer point, or
- the fuel measurement point has a clean, dry fuel source at a facility, such as a gas plant.

If the current metering installation does not meet the grandfathering requirement, operators must make any necessary changes required to bring the installation into compliance with this section within one year of **February 2, 2009.**

4.3.4.2 Chart Operations

Field operation personnel **must** ensure that

- the meter location is properly identified on the chart;
- the chart is correctly dated;
- the on and off chart times are recorded on the chart to the nearest quarter hour if not actual;
- the correct orifice plate and line size are recorded on the chart;
- the time (to the nearest quarter hour) of any orifice plate change is indicated on the chart, along with the new orifice plate size;
- it is noted on the charts if the differential, pressure, or temperature range of the recorder has been changed or if they are different from the ranges printed on the chart;
- a copy of the chart calibration report is kept on site or readily available for on-site inspection if it is a manned facility;
- the flowing gas temperature is recorded on the chart in accordance with Table 4.2 above;

- proper chart reading instructions are provided when the pen fails to record because of sensing line freezing, clock stoppage, pens out of ink, overlapping traces, or other reasons (draw in the estimated traces, request to read as average flow for the missing period, or provide estimate of the differential and static); and

- any data or traces that require correction must not be covered over or obscured by any means.

Field operation personnel **should** ensure that

- a notation is made on the chart with regard to whether or not the meter is set up for atmospheric pressure (for square root charts);

- the accuracy of the meter clock speed is checked and the chart reader is instructed accordingly of any deviations;

- the differential pen is zeroed once per chart cycle;

- differential pen recordings are at 33% or more within the chart range whenever possible;

- static pen recordings are at 20% or more within the chart range whenever possible;

- when there is a painted differential band, instructions are provided as to where it should be read; there are various ways to read a painted chart:

  - if the differential pen normally records at the top of the painted band but spikes quickly down and up during separator dump cycles, it is reasonable to read the differential near the top of the band (or vice versa);

  - if the differential pen is in constant up and down motion, it is reasonable to read the differential at the root mean square (RMS) of the band or in a sine wave motion alternating between the top and bottom of the painted area;

- the pen trace colours conform to the industry-accepted practice (RED for differential, BLUE for static, and GREEN for temperature); however, any colour may be used, provided the colour used is documented.

4.3.4.3 ERCB Site-Specific Requests

If an inspection of a measurement device or of procedures reveals unsatisfactory conditions that significantly reduce measurement accuracy, the ERCB inspector or auditor will direct in writing that the licensee implement changes to improve measurement accuracy, and this direction will become a condition of operation for that facility or facilities. Examples of conditions applicable to orifice chart recorders are as follows:

- thick pen traces that will cause excessive error when reading the traces

- painting traces

- differential or static pens recording too low on the chart—in some cases, this cannot be avoided because of low flow rate, high shut-in pressure, and equipment or operating pressure range limitations.

4.3.4.4 Chart Reading

The chart integrator/planimeter operator **must** ensure the following:

- Visible gaps between the integrator/planimeter traces and chart traces are minimized.

- The counter is read correctly.

- The integrator is calibrated periodically and after each change of pens.

- The correct integrator or square root planimeter constants are noted.
- The correct integrator setback is recorded.
- The correct coefficient, using all of the required factors, is recorded.

4.3.4.5 Alternative Chart Reading Technology

The base requirements for alternative methods developed to read orifice meter charts, other than conventional manual methods (planimeters, integrators), is as follows. An example of such technology is the use of digital scanning technology to scan and store an image of the chart and the use of computer programs to read and interpret the digital image of the chart and the pen traces.

The use of alternative technologies to read charts does not require prior approval of the ERCB, but the licensee using any new technology must be able to demonstrate that the following requirements are met:

- The equipment and/or procedures used to read the chart must not alter or destroy the chart such that it cannot subsequently be read using conventional equipment and/or procedures.

- The accuracy and repeatability of the new equipment and/or procedures must be equal to or better than conventional equipment and/or procedures.

The following requirements are specific to the use of digital scanning technology for reading charts:

1) The original chart must be retained for at least 12 months (18 months for gas production associated with heavy oil or crude bitumen), or alternatively the licensee may choose the following procedure for audit trail:

   a) An original scanned image of the chart (both front and back) must be stored so that it cannot be changed. If the chart back is blank, the back does not need to be scanned provided there is a statement entered in the record to that effect. There must be a method to confirm that a set of front and back scans belong to the same chart if scanned and stored. No alteration or editing of the original scanned image is allowed.

   b) At least two separate electronic copies of the scanned images must be retained and one copy must be stored off site at a different physical address/location for the applicable required period.

   Note that although the ERCB accepts the above electronic submission for audits, other jurisdiction might not. Therefore the original chart should be kept for other jurisdictional audits.

2) Editing or alterations may only be made to a copy of the original scanned image of the chart. If the edited version is used for accounting purposes, the edited or altered image must be stored for the applicable required period and in the same manner as in item 1 above.

3) An image of the chart showing how the chart pen traces were read or interpreted must be stored for the applicable required period and in the same manner as in item 1 above.

4) The requirements and recommendations in Section 4.3.4 of this directive must be adhered to. If there are any changes or additions to those requirements and recommendations specific to chart scanning, these must be documented and made available for instructing chart analysts. An additional requirement specific to chart scanning is as follows:

- When a differential pen is not zeroed correctly, the zero line must be adjusted to the correct position if it is obvious on the chart (such as when the zeroing was out when changing charts but the pen was not adjusted) and/or as documented by the operator. Other situations will require the judgement of the chart analyst and confirmation from the facility operator. Any zero adjustment must only reposition the zero line and must maintain the entire span of the pen. (The distance between the actual zero and the pen trace must not be altered.)

5) For ERCB inspection/audit purposes, the licensee must upon request

- submit any original paper charts or the scanned original images or make them available for on-line viewing, and

- submit all edited images or make them available for on-line viewing.

Note that the software used to open the scanned images should be readily and freely available on the market. In case there is any specific/proprietary image reader software required to view the scanned and stored chart images, it must also be submitted.

6) Upon request, the repeatability of the scanning technology must be demonstrated by performing three consecutive scans with a rotation of the chart image of about 120° before each scan and integrations of the same chart image. The calculated volumes from each reading must be within ± 0.5% of the average of the three scans and integrations.

7) The ERCB may check the accuracy of the chart-reading technology and volume calculations by providing charts with known calculated volumes. The volumes determined by the chart reading technology must be within ± 0.5% of the ERCB's known values.

4.3.5 Exceptions

4.3.5.1 Gas in Solution with Oil Volumes under Pressure

In some cases a gas volume must be determined where the gas is dissolved in an oil volume under pressure, and there is no opportunity to measure the gas volume prior to it being commingled with other gas volumes. In that case, the gas volume may be determined by estimation, regardless of its daily volume rate. An example of such a gas volume is the gas held in solution with oil volumes leaving a test separator at an oil proration battery, where the test oil volumes are combined with production from other wells downstream of the test separator. The purpose of estimating the gas in solution is to determine the total gas produced by a well during a production test, since the gas volume measured by the test gas meter will not include the gas that is still in solution with the test oil volume.

A single gas-in-solution (GIS) factor may be determined and used to estimate the gas volume held in solution with the oil stream for each oil stream where the production sources (producing formation) are the same and test separator operating conditions are similar. Additional GIS factors are required for wells in the battery that produce from different formations and where other test separators operate at different pressure and/or temperature conditions. Licensees should also consider determining seasonal GIS factors where ambient

temperature differences may significantly affect the factors or when operating conditions change significantly.

The GIS factor may be determined by one of the following applicable tests/procedures:

- A 24-hour test may be conducted such that the production from a well (or group of wells) is directed through the test and group separation/treating equipment, with all other wells shut in or directed around the equipment. The total volume of gas released from the oil after it leaves the test separator must be measured; this volume divided by the stock tank volume of oil determined at the test separator provides a GIS factor.

- A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a pressure-volume-temperature (PVT) analysis can be conducted. The analysis must be based on the actual pressure and temperature conditions that the oil sample would be subjected to downstream of the sample point, including multiple stage flashing. The GIS factor is calculated based on the volume of gas released from the sample and the volume of oil remaining at the end of the analysis procedure.

- A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a compositional analysis can be conducted. A computer simulation program may be used to determine the GIS factor based on the compositional analysis.

- A "rule of thumb" estimate (0.0257 m^3 of gas/m^3 of oil/kPa of pressure drop) may be used as the GIS factor for conventional light-medium oil production until a more accurate, specific GIS factor is determined. It may be used on a continuous basis, without the need for determining a more accurate GIS factor, if well oil production rates do not exceed 2 m^3/d or if all battery gas production is vented or flared.

- Other methods listed in the Canadian Association of Petroleum Producers (CAPP) *Guide for Estimation of Flaring and Venting Volumes from Upstream Oil and Gas Facilities* may be used.

4.3.5.2 Gas Produced in Association with Conventional Oil Well and Gas Well Production

If a gas stream volume associated with conventional oil well or gas well production does not exceed $0.5 \ 10^3$ m^3 per day at any given measurement/disposition point, the volume may be determined by estimation instead of measurement. **No specific approval is required, but the operator must keep the estimation/testing documentation for ERCB audit.** Examples of the gas streams that may be estimated if the daily volume limitation is not exceeded include well test gas, battery group gas, single-well battery gas, fuel gas used on a per site basis, and oil/condensate tank vented gas. A gas stream that must be measured regardless of daily volume is dilution gas added to an acid gas stream to ensure complete combustion (due to the importance of accurately determining those volumes).

Initial qualification of gas streams where volumes may be estimated can be based on existing historical data or determined by conducting one of the applicable tests/procedures in Section 4.3.5.4. Qualifying gas volumes may be estimated by using a gas-oil-ratio (GOR) factor if gas volume estimates will vary in conjunction with oil volumes or by using an hourly rate if gas volumes are not dependent upon oil volumes. These factors must be updated annually to confirm continuing eligibility for estimation and to update the factors used to estimate gas volumes. The factors must also be updated immediately following any operational changes

that could cause the factors to change. Licensees should also consider determining seasonal GOR factors if ambient temperature differences may significantly affect the factors. Updated factors may be determined by one of the applicable tests/procedures described below.

4.3.5.3 Gas Produced in Association with Heavy Crude Oil/Crude Bitumen Production

See Section 12.3.2 for details.

4.3.5.4 Methods for Determining Factors/Rates Used in Estimating Gas Volumes

If gas volumes will be estimated using a GOR:

- A 24-hour test may be conducted such that all the applicable gas and oil volumes produced during the test are measured (including vented gas). The gas volume is to be divided by the oil volume to result in the GOR factor.

- A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a PVT analysis can be conducted. The analysis must be based on the actual pressure and temperature conditions the oil sample would be subjected to downstream of the sample point. The GOR factor will be calculated based on the volume of gas released from the sample and the volume of oil remaining at the end of the analysis procedure.

- A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a compositional analysis can be conducted. A computer simulation program may be used to determine the GOR based on the compositional analysis.

- Other methods listed under the Canadian Association of Petroleum Producers (CAPP) *Guide for Estimation of Flaring and Venting Volumes from Upstream Oil and Gas Facilities* may be used

If gas volumes will be estimated using an hourly rate:

- A meter may be used to measure the gas stream for a minimum of one hour. The gas volume measured during this test may be used to determine the hourly rate that will be used to estimate gas volumes.

- If applicable, such as for fuel gas volumes, the hourly rate may be determined based on the equipment manufacturer's stated gas consumption rates and the actual operating conditions.

Example Calculations for Estimating Gas Volumes Using GOR and GIS Factors

Determination of Total Produced Gas for a Single-Well Oil Battery

Figure 4.4 depicts a single-well battery where a three-phase separator is used to separate oil, gas, and water production from a well. The oil in the separator is under pressure until it is directed to the storage tank, which is at atmospheric pressure (zero kPa gauge). When the oil pressure drops at the tank, the GIS within the oil will be released. The gas leaving the separator in this example is measured, while the GIS released at the tank is estimated using a GOR factor. Total gas production from the well is determined by adding the measured gas and the GIS released at the oil storage tank.

If a single-well battery uses a two-phase separator, the procedure for determining total gas production is the same as for a three-phase separator.

If the gas production rate meets the qualifying criteria for estimation and all production from the well produces directly to a tank without using a separator, the total gas production may be determined by using only a GOR factor.
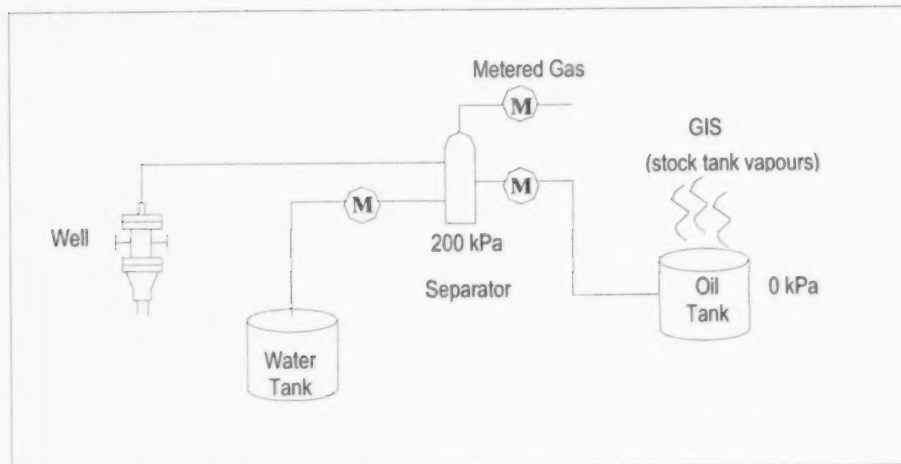
Figure 4.4. Single-well oil battery example

Sample Calculation: Total Gas Volume at a Single-Well Battery (Figure 4.4)

Monthly well data (hypothetical) given for this example:

Gas meter volume = 96.3 10^3 m^3 (from chart readings)

Oil meter volume = 643.3 m^3 (from meter or tank gauging)

Pressure drop = 200 kPa

GOR factor = 6.37 m^3 gas/m^3 oil or 0.03185 m^3 gas/m^3 oil/kPa pressure drop (determined using a method other than the "rule of thumb")

Step 1: Calculate GIS volume:

6.37 m^3/m^3 x 643.3 m^3 = 4097.8 m^3 = 4.10 10^3 m^3
or
0.03185 m^3/m^3/kPa x 643.3 m^3 x 200 kPa = 4097.8 m^3 = 4.10 10^3 m^3

Step 2: Calculate the total battery gas production for the month:

96.3 10^3 m^3 + 4.1 10^3 m^3 = 100.4 10^3 m^3

Note that total reported battery gas production is to be rounded to one decimal place.

Determination of Total Produced Gas for an Oil Proration Battery

Figure 4.5 depicts a multiwell oil proration battery where production testing of individual wells is done by directing individual well production through a test separator at the main battery site or through a test separator at a satellite facility located away from the main battery site.

In this example, the oil, gas, and water leaving the test separator at the satellite are recombined with the satellite group production and directed to the group separation and measurement equipment at the main battery site. The oil and water leaving the test separator at the main battery site are recombined with the battery group production, but the gas leaving the test separator recombines with the group gas downstream of the group gas measurement point. The oil in the group separator is under pressure until it is directed to the storage tank, which is at atmospheric pressure (zero kPa gauge). When the oil pressure drops at the tank, the GIS with the oil will be released.

The total gas production at the battery will be the sum of all the measured test gas at the battery site, the measured group gas at the battery, and the GIS released at the oil storage tank.

Trucked oil volumes received at the battery must not be included with the total battery oil volume when determining the GIS released at the oil storage tank.

At some facilities a vapour recovery unit (VRU) may be installed to collect any GIS that may be released at the oil storage tank. If the VRU is equipped with a meter or the recovered gas is directed through the group gas meter, a GIS calculation will not be required because the measured VRU gas will either be added to or included in the other measured gas volumes.
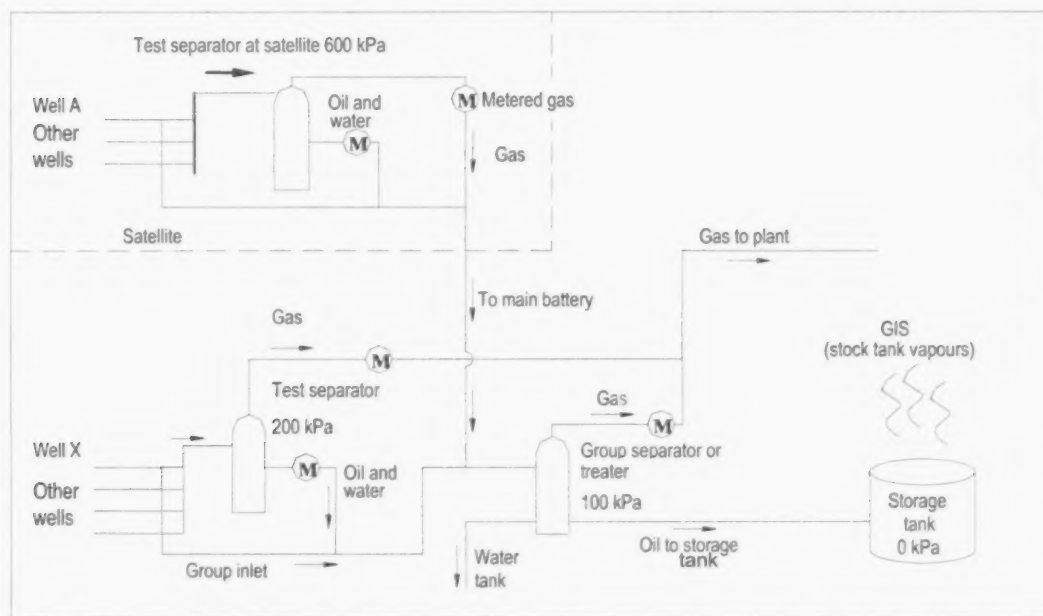
Figure 4.5. Multiwell oil battery example

Sample Calculation: Total Gas Production at the Oil Proration Battery (Figure 4.5)

Monthly battery data (hypothetical) given for this example:

Oil production at the proration battery = 745.0 m^3 for the month (from meter and/or tank gauging)

Total test gas measured at the battery site = 30.0 10^3 m^3 (from chart readings)

Measured group gas production = 67.4 10^3 m^3 (from chart readings)

Pressure drop from the group vessel to oil storage tank = 100 kPa

GOR factor = 3.99 m³ gas /m³ oil or 0.0399 m³/m³/kPa (determined using a method other than the "rule of thumb")

Step 1: Calculate the GIS volume:

3.99 m³/m³ x 745.0 m³ = 2972.6 m³ = 2.97 10³ m³

or

0.0399 m³/m³/kPa x 745 m³ x 100 kPa = 2972.6 m³ = 2.97 10³ m³

Step 2: Calculate the total produced gas volume for the battery:

67.4 10³ m³ + 30.0 10³ m³ + 2.97 10³ m³ = 100.4 10³ m³

Note that total reported battery gas production is to be rounded to one decimal place.

Determination of Individual Well Test Gas for an Oil Proration Battery

Figure 4.5 depicts a multiwell oil proration battery where production testing of individual wells is done by directing individual well production through a test separator at the main battery site or through a test separator at a satellite facility located away from the main battery site. In either case, the oil leaving the test separator is under pressure and will be subjected to two stages of pressure drop—one at the group separator and one at the storage tank. The total gas produced by a well during a test will be the sum of the gas measured as it leaves the test separator and the GIS that will evolve from the test oil volume after leaving the test separator. In the example, the test separators at the battery and satellite operate at significantly different pressures, and the oil leaving the test separator at the satellite will contain more GIS than the oil leaving the test separator at the battery.

Sample Calculation: Test Gas Production for Wells in the Satellite (Figure 4.5)

Satellite test data (hypothetical) given for this example for well "A":

Measured test oil = 7.22 m³ (from oil meter)

Measured test gas = 1.27 10³ m³ (from chart readings)

GIS factor = 25.62 m³ gas/m³ oil or 0.0427 m³ gas/m³ oil/kPa pressure drop (combined GIS for both stages of pressure drop from test pressure at 600 kPa to group pressure at 100 kPa to oil storage tank pressure at atmospheric pressure or zero kPa gauge, determined using a method other than the "rule of thumb")

Step 1: Calculate the GIS volume:

0.0427 m³/m³/kPa x 7.22 m³ x 600 kPa = 185.0 m³ = 0.19 10³ m³

or

25.62 m³/m³ x 7.22 m³ = 185.0 m³ = 0.19 10³ m³

Step 2: Calculate the total test gas produced for well "A" for this test:

1.27 10³ m³ + 0.19 10³ m³ = 1.46 10³ m³

Note that test gas volumes must be determined to two decimal places (in 10³ m³).

Sample Calculation: Test Gas Production for Wells in the Battery (Figure 4.5)

Battery test data (hypothetical) given for this example for well "X":

Measured test oil = 3.85 m^3 (from oil meter)

Measured test gas = 2.33 10^3 m^3 (from chart readings)

GIS factor = 7.90 m^3 gas/m^3 oil or 0.0395 m^3 gas/m^3 oil/kPa pressure drop (combined GIS for both stages of pressure drop from test pressure at 200 kPa to group pressure at 100 kPa to oil storage tank pressure at atmospheric pressure or zero kPa gauge, determined using a method other than the "rule of thumb")

Step 1: Calculate the GIS volume:

0.0395 m^3/m^3/kPa x 3.85 m^3 x 200 kPa = 30.4 m^3 = 0.03 10^3 m^3
or

7.90 m^3/m^3 x 3.85 m^3 = 30.4 m^3 = 0.03 10^3 m^3

Step 2: Calculate the total test gas produced for well "X" for this test:

2.33 10^3 m^3 + 0.03 10^3 m^3 = 2.36 10^3 m^3

Note that test gas volumes must be determined to two decimal places (in 10^3 m^3).

4.4   References

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5 Site-specific Deviation from Base Requirements

Section 1: Standards of Accuracy states that a licensee may deviate from the ERCB's minimum measurement, accounting, and reporting requirements without specific approval if no royalty, equity, or reservoir engineering concerns are associated with the volumes being measured and the licensee is able to demonstrate that the alternative measurement equipment and/or procedures will provide measurement accuracy within the applicable uncertainties. This section describes situations where a licensee may deviate from the minimum requirements without ERCB approval, provided that specific criteria are met. Licensees may also apply for approval to deviate from the minimum requirements if the specific criteria are not met; this section indicates what information must be included in such an application. If these exceptions or approvals are in use, ERCB inspectors and auditors will review the licensee's records for demonstrated compliance with the criteria specified in this section or in the applicable approval.

Specialized Terminology Defined

Common Ownership	All wells in a battery belong to the same working interest participant, or if there is more than one working interest participant, each working interest participant has the same percentage interest in each well in the battery.
Common Crown or Freehold Royalty	When all the wells in a battery are produced under Crown mineral leases, the Crown receives the same royalty rate for each well, or when under leases granted by one Freehold mineral holder, the Freehold mineral holder receives the same royalty rate for each well. If there is more than one Freehold mineral holder for the wells in a battery, the total royalty rate for each well is the same.
Measured Gas Source(s)	These are single-phase measured gas source(s) downstream of separation and removal of liquids. This also includes the gas equivalent volume (GEV) of measured condensate if the condensate is recombined with the gas downstream of the separator.
Measured Oil	Oil is measured using equipment and/or procedures meeting delivery point measurement uncertainty limits. For emulsion, the delivery point measurement uncertainty limits apply to the total volume determination only.

5.1 Site-specific Exceptions

Deviation from base measurement, accounting, and reporting requirements is allowed without submission of an application to the ERCB, provided that all the initial qualifying criteria listed under the subsequent "Exception" sections are met.

Initial Qualifying Criteria—These criteria (detailed in subsequent sections) must be met to qualify for the exception. If the initial qualifying criteria have been met and the exception is implemented, it may remain in place indefinitely, as long as it does not meet any of the revocation clauses and no physical additions to the facility are made (e.g., new wells or zones). If additions or changes are made to the battery or facility, the initial qualifying criteria must be met for all the wells or zones added to the battery or facility for the exception to remain in place. If the licensee anticipates that physical additions may not meet the initial

qualifying criteria, the licensee may reconfigure the facility to meet base measurement, accounting, and reporting requirements or submit an application for site-specific approval of deviation from the base requirements. Approval must be in place prior to implementation. Submission of an application does not guarantee that an approval will be granted.

Documentation Requirement—The licensee must retain the data and documentation to support the initial qualifying criteria and the last three testing records (if applicable) for as long as the exception is in place. The ERCB may revoke an exception if an audit or inspection reveals a lack of adequate supporting data or documentation. If the licensee cannot provide documentation requested for ERCB audit within 30 days, the licensee will be required to meet applicable ERCB base measurement requirements immediately. Alternatively, at the ERCB's discretion, the licensee may propose a plan to comply with the ERCB exception requirements within an ERCB-approved time period.

5.2 Site-specific Approval Applications

If the exception criteria cannot be met or if a specific situation is not covered in this section, the licensee may be allowed to deviate from base measurement, accounting, and reporting requirements upon approval of an application submitted to the ERCB Production Operations Section.

Approvals will remain in place indefinitely, including after transfer of the facility to another licensee, provided that conditions specified in the approval are met. If an ERCB audit or inspection finds that approval conditions are not being met, the approval may be revoked and the licensee may be required to meet applicable base requirements immediately, or other appropriate requirements may be specified.

If a licensee anticipates that proposed changes to the facility may not meet the approval conditions, the licensee may reconfigure the facility to meet base measurement, accounting, and reporting requirements or submit a new application for site-specific approval of deviation from the base requirements. Approval must be in place prior to implementation. Submission of an application does not guarantee that an approval will be granted.

The following information is required for all applications for site-specific deviation from base requirements. Other specific information that may be required is described in the following appropriate sections.

1) Well and/or facility list, including

- battery code and locations
- well locations (UID)
- licence number(s)
- respective pool/zone designations and unique identifier for each zone
- indication as to unit or non-unit operation, if applicable
- royalty status (Freehold/Crown, new/old, etc.)
- equity (ownership) issues, if any
- latest 6 months' gas, oil/condensate, and water flow rates (or expected flow rates for new wells)
- up-to-date measurement schematic(s) for the existing system(s) and the proposed new gas or oil source(s), including all tie-in locations, if applicable

- facility plot plan for the existing system and the proposed new gas or oil source(s), if applicable

2) Justifications for deviation from measurement requirements (e.g., economics, minimal impact on measurement accuracy)

5.3 Chart Cycles Extended Beyond the Required Time Period

Chart cycle is the time required for a circular chart to complete one 360° revolution. An extension of the required chart cycle time may be applicable under the following scenarios:

- the gas well orifice meter chart cycle is greater than 8 days;
- the single-well oil battery orifice gas meter chart cycle is greater than 24 hours; or
- the Class 3 and 4 oil well test orifice gas meter chart cycle is greater than 8 days.

Mixing of wells with EFM systems and wells using extended cycle paper charts within the same battery is allowed without approval from the ERCB.

Group, sales, or delivery point meters and Class 1 and 2 oil well test gas meters do not qualify for exception for chart cycle extension, and approvals for extension of the chart cycle for those meters will not normally be granted.

5.3.1 Exceptions

Orifice meter gas chart cycles may be extended without ERCB site-specific approval if all the initial qualifying criteria below are met and no application is required.

Initial Qualifying Criteria

1) In the case of gas well measurement, all wells in the multiwell battery are gas wells. A single-well battery does not qualify for this exception on its own; the entire group battery or gas gathering system must be considered.

2) In the case of oil well measurement, all wells in the battery are oil wells, and the well produces either to a single-well battery or to a multiwell oil group where each well has its own separation and measurement equipment.

3) All wells are subjected to the same type of measurement (all well production is separated and all components are measured, or all well production is subject to effluent measurement) and the same chart cycle.

4) All wells flowing to the battery have common ownership and either common Crown or Freehold royalty, or

- if there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received;

- if there are no common Crown or Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalty involved, the licensee must apply to the ERCB for approval.

5) The monthly average volumetric gas flow rate for each well is $\leq 16.9 \ 10^3 \ m^3/d$ (including the gas equivalent of condensate in the case of gas well measurement).

6) The differential pen records at 33% or more within the chart range, and the static pressure pen should record at 20% or more within the chart range (if possible). Painted traces must not exceed 4% of the differential pressure or static pressure range. Painting occurs when there are quick up and down movements of the pen, so that there is no visible separation between the up and down traces for a period of time.

7) Temperature must be recorded at a minimum of once per week, and if that is not possible, continuous temperature measurement (temperature pen) is required.

8) The wells that are within the same battery of the extended chart cycle wells and are designed for and/or operate on on/off flows (e.g., plunger lifts, pump-off controls, intermittent timers) must be measured using EFM. In addition, an extended chart cycle with EFM is allowed.

Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) Oil well/battery gas is added to a gas battery.

2) There is mixed measurement within the battery other than with EFM.

3) The oil well is not produced either to a single-well battery or to a multiwell oil group where each well has its own separation and measurement equipment.

4) The working interest participants for any well flowing to the battery have changed and a new working interest participant objects to the exception.

5) Any well within the battery has exceeded the $16.9 \ 10^3 \ m^3/d$ monthly average actual gas production rate (including gas equivalent of condensate for gas wells).

6) Painted traces for any well exceeded 4% of the differential pressure range or the static pressure range.

7) A new well with on/off flows is added to an effluent proration battery, or one or more of the existing wells has been modified to operate on on/off flows but EFM is not used.

Base measurement requirements must be reinstated if the exception is revoked due to any of the above.

5.3.2 Applications

The following information must be submitted with an application to extend orifice meter gas chart cycles if the above criteria are not met:

1) all of the information listed in Section 5.2: Site-specific Approval Applications;

2) if there are no common Crown or Freehold royalties or common ownership, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold royalty holders and working interest participants, with no resulting objection received;

3) a discussion of the impact on measurement accuracy of intermingling base chart cycles and extended chart cycles in a common battery, and how it may relate to concerns about working interest equity and/or royalty considerations;

4) a minimum of two current, consecutive, representative gas charts—Additionally, the licensee has the option to run the charts on the proposed chart cycle to gather test data for submission and then revert back to the required chart cycle after a maximum test period of 31 days. The original copies of any such charts created must be submitted with the application. The trial run must be clearly identified on the charts.

5.3.3 Considerations for Site-specific Approval

1) Differential and static pressures are stable, with essentially uninterrupted flow:

- On/off flow as designed, including plunger lifts, pump-off controls, intermittent timers, etc., that cause painting or spiking, do not normally qualify for chart cycle extension.

- The effects of painting are minimized. The amount of painting that is acceptable is decided case by case.

- The differential pen should record at 33% or more within the chart range and the static pressure pen should record at 20% or more within the chart range (if possible).

2) There are minimal equity and royalty concerns.

3) Reservoir engineering concerns: The concern for well measurement accuracy declines, from a reservoir perspective, as the pool depletes. The applicant should provide its assessment/opinion, but the ERCB has to decide on a case-by-case basis if the concerns are relevant.

4) All gas meters producing into the same group measurement point use the same chart cycle, so that they are subject to the same type of error.

5.4 Gas Proration Outside Southeastern Alberta Shallow Gas Zones/Area

For wells outside the boundary of and/or producing from zones other than those approved for the Southeastern Alberta Shallow Gas Zones/Area (see 7.2), it may be acceptable to use a proration system for gas well production instead of having measurement for every well. If a proration system is implemented, all wells in the battery must be subject to the proration system.

5.4.1 Exceptions

Gas wells may be produced without individual well measurement and be connected to a proration battery without ERCB site-specific approval if all the initial qualifying criteria below are met and no application is required.

Initial Qualifying Criteria

1) All wells are classified as gas wells.

2) All wells flowing to the battery have common ownership and either common Crown or Freehold royalty, or

- if there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received;

- if there are no common Crown or Freehold royalties and there are only Freehold royalties involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to the ERCB for approval.

3) The licensee has discussed and addressed reservoir engineering issues with its own reservoir engineering staff or external knowledgeable personnel to ensure minimal reservoir engineering concerns and has documented the results for audit.

4) Total liquid production at each well in the battery is ≤ 2 m^3/d based on the monthly average flow rates recorded during the six months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.

5) The maximum average daily well gas flow rate of all wells in the battery is $\leq 10.0 \ 10^3$ m^3/d (including gas equivalent volume of condensate), with the highest daily well flow rate $\leq 16.9 \ 10^3$ m^3 (including gas equivalent volume of condensate and except as allowed below). If an existing battery with measured gas well production is being converted to a proration battery, qualifying flow rates must be based on the monthly average flow rates recorded during the six consecutive months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.

6) Periodic well tests are conducted under normal operating conditions to determine hourly flow rates that will be used to estimate monthly well production based on monthly well operating hours. The well tests are conducted for a minimum of 12 hours, and all gas, condensate, and water volumes are separated and measured during the test. For gas wells with minimal water production (≤ 0.01 m^3 water/10^3 m^3 gas) and no condensate or oil, the testing duration must be sufficient to clearly establish stabilized flow rates and single-phase testing is allowed,

7) Following the commencement of production at the proration battery, all wells are tested within the first month, then again within six months, and then annually after that. New wells added to the battery at some future date must be tested within the first month of production, then again within six months, and then annually after that.

8) For new wells tying into a gas proration battery and that will be producing more than 16.9 10^3 m^3/d but that are expected to drop below 16.9 10^3 m^3/d within six months, every well must be tested monthly for the first six months with a separator, or until the production rate has stabilized, and annually thereafter. If the gas production rate for any of the wells is more than 16.9 10^3 m^3/d after six months of production or the liquid production rate is higher than 2 m^3/d, a separator must be installed to continuously separate and measure the well production, and the measurement-by-difference rules in Section 5.5 apply in this case.

9) For coalbed methane (CBM) wells and wells producing from above the base of groundwater protection each with water production ≤ 0.01 m^3 water/10^3 m^3 gas and no

condensate, if at any time more than 5.0 m³/month of net water production is realized at the group measurement point, the operator must investigate the source of the water production by retesting and using at least a two-phase separator at the suspected gas well(s) within 30 days and then prorate the water production accordingly.

10) The flow rates established from the well tests are used to determine estimated monthly well production from the date of the test until the date of the next test, with the exception that the test conducted during the first month of production is also used to estimate the wells' production for the producing days prior to the test. The total measured group gas and liquid production are prorated to the wells, based on each well's estimated production, to determine the actual well production.

Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) An oil well is added to the battery, or one or more of the existing gas wells has been reclassified as an oil well.

2) The maximum average daily flow rate of all wells in the battery for any month exceeded 10.0 10³ m³/d, or the highest single well flow rate exceeded 16.9 10³ m³/d except as allowed above.

3) Total liquid volume exceeded 2 m³/d during a 24-hour test period or prorated to 24 hours if the test period is not 24 hours.

4) A new well has been added to the proration battery with a daily flow rate over 16.9 10³ m³ except as allowed above or whose additional volume will cause the average daily well gas flow rate of all wells in the battery to exceed 10.0 10³ m³/d.

5) Wells within the proration battery or new wells added to the battery were not tested as required.

6) The gas proration methodology in item 10 under Initial Qualifying Criteria above was not followed.

Base measurement requirements must be reinstated if the exception is revoked due to any of the above.

5.4.2 Applications

The following information must be submitted with an application to use a proration system, instead of individual gas well measurement, to determine gas well production if the above criteria are not met:

1) all of the information listed in Section 5.2: Site-specific Approval Applications;

2) a discussion of the stage of depletion for pools involved and the impact of any reduction in well measurement accuracy that may result from gas proration as it relates to reservoir engineering data needs—discussion of this matter by the licensee with its own reservoir engineering staff or knowledgeable external personnel is required and must be addressed in the application;

3) a clear explanation and flow diagram of proposed well and group measurement devices and locations, the proposed accounting and reporting procedures, and the proposed method and frequency of testing;

4) if there is no common Crown or Freehold royalties or common ownership, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold royalty holders and working interest participants, with no resulting objection received.

5.4.3   Considerations for Site-specific Approval

1) All wells must be classified as gas wells.

2) There are minimal equity, royalty, and reservoir engineering concerns.

3) All wells should have similar flow rates.

4) Economic considerations: Would implementation of a proration system reduce costs enough to significantly extend operations? Have other options been considered?

5) Total liquid production at each well in the battery should be $\leq 2 \ m^3/d$ based on the monthly average flow rates recorded during the six months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.

5.5   Measurement by Difference

Measurement by difference is defined as any situation where an unmeasured volume is determined by taking the difference between two or more measured volumes. It results in the unmeasured volume absorbing all the measurement error associated with the measured volumes. In the case of a proration battery (effluent measurement, or periodic testing without continuous measurement), new gas or oil source errors may be difficult to detect because the proration testing errors in the original system can hide the new source errors. Despite these concerns, a properly designed and operated measurement system can minimize the risk and attain reasonable accuracy, provided that the measured source gas or oil rates are a small proportion of the total system delivery rates. Measurement by difference is not allowed for "gas group batteries," "single-well batteries," or "sales points" unless special approval is obtained from the ERCB.

Gas Measurement by Difference

For gas proration batteries, measurement by difference can include but is not limited to the following situations. (Note that all schematics below are examples only; some systems may be configured differently.)

Measured gas source(s) (other than from the designated SE Alberta Shallow Gas Zones/Area) delivering into a SE Alberta Shallow Gas proration battery (Figure 5.1):
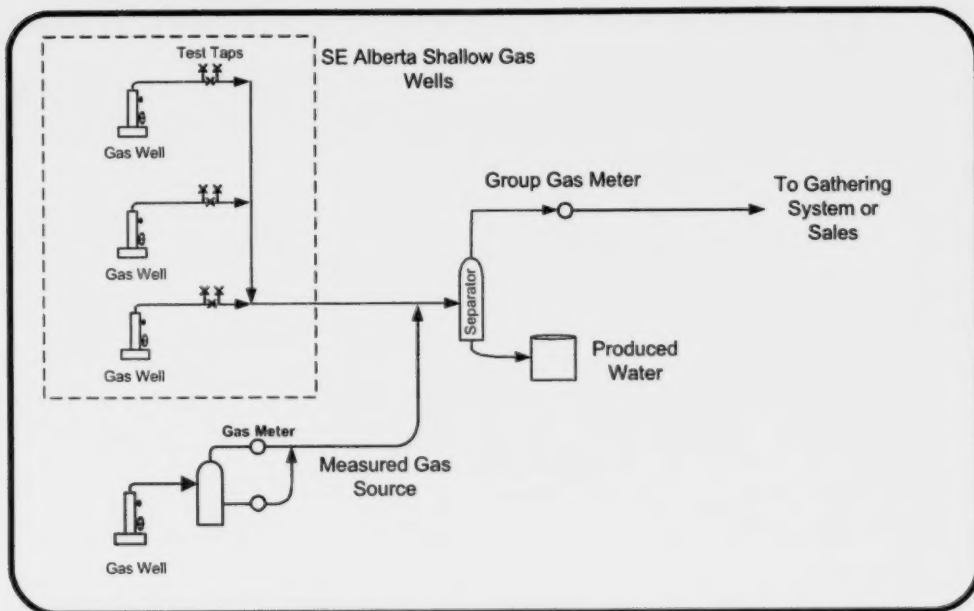
Figure 5.1

- Measured gas source(s) delivering into a gas proration battery outside SE Alberta Shallow Gas Zones/Area (Figure 5.2):
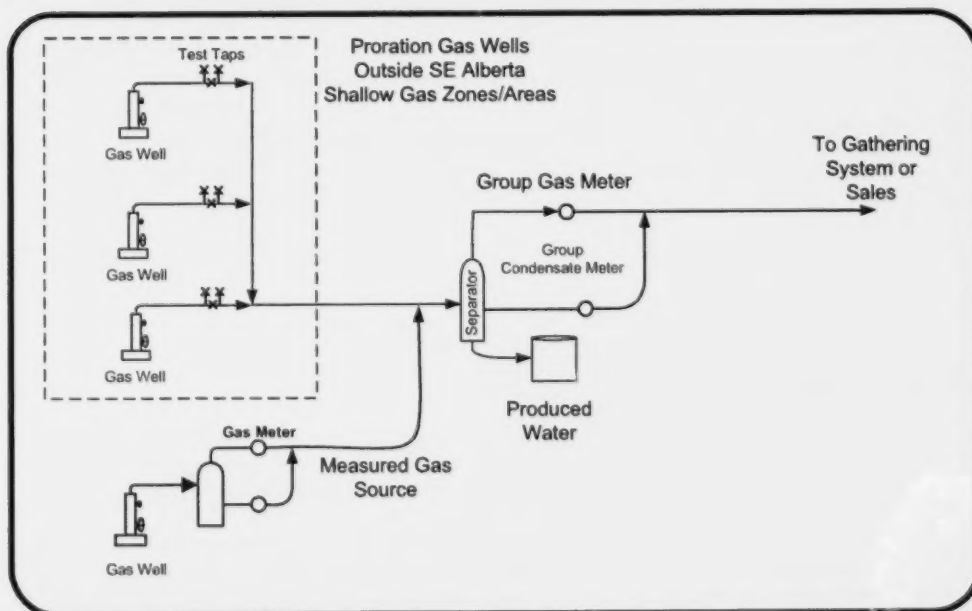
Figure 5.2

- Measured gas source(s) delivering into an effluent (wet) measurement gas proration battery (Figure 5.3):
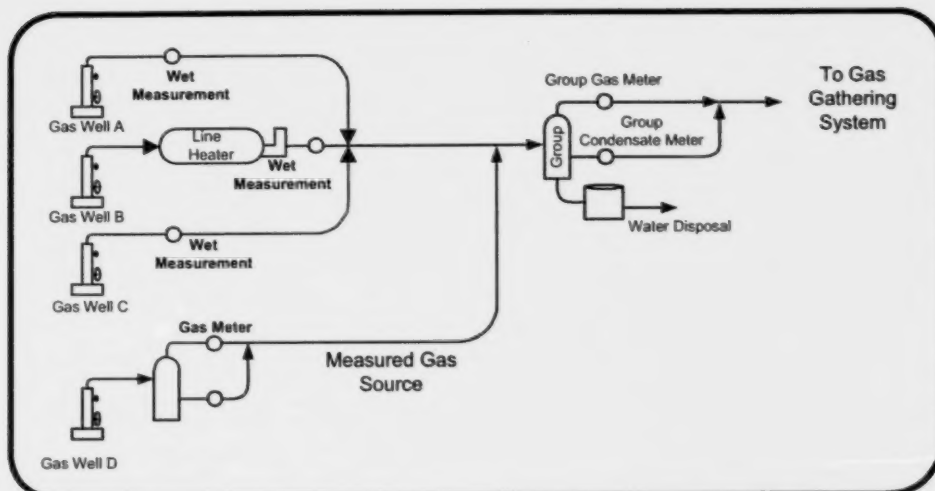
Figure 5.3

- Measured gas source(s) delivering into an oil proration battery (Figure 5.4):
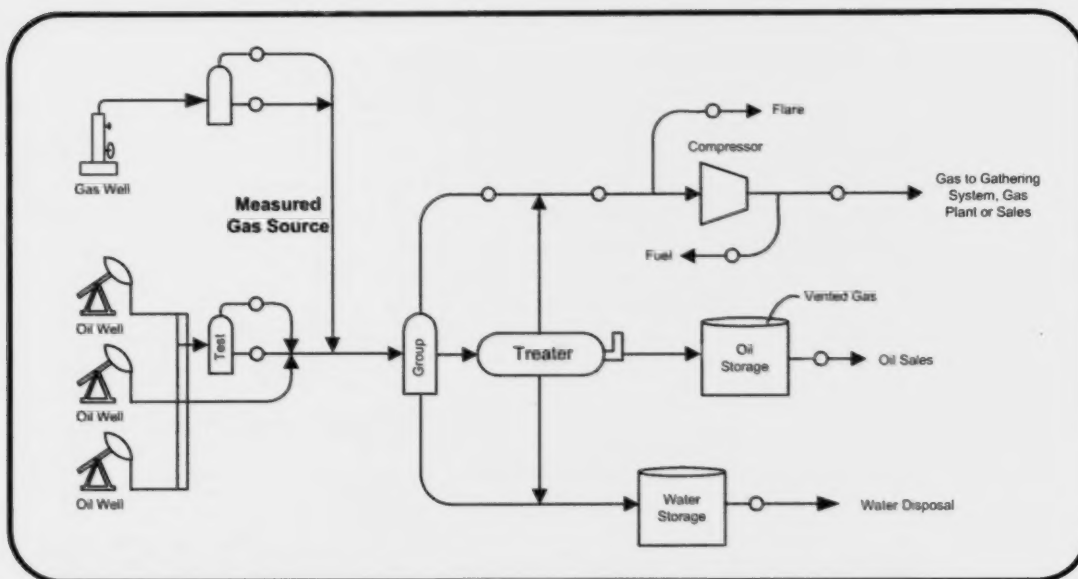
Figure 5.4

- Measured oil facility gas delivering into a gas proration battery (Figure 5.5):
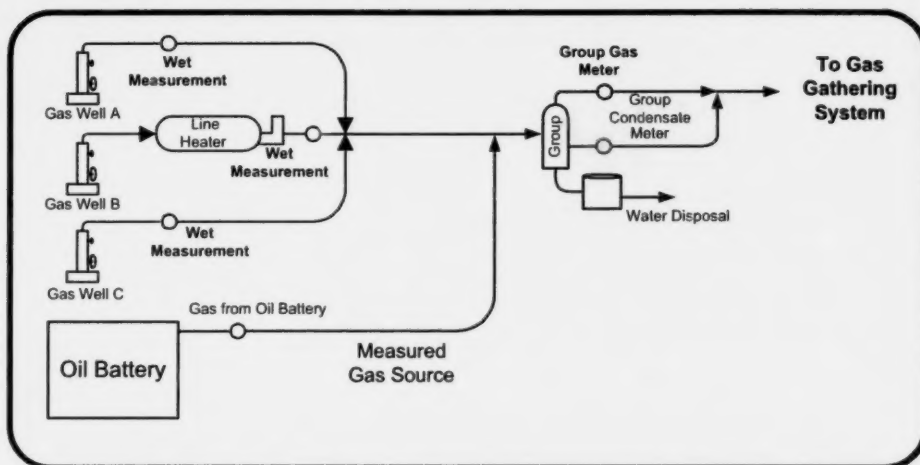
Figure 5.5

If any measured gas source will be tied in to a gas proration battery:

1) The gas and liquids from all tied-in gas sources must be separately and continuously measured.

2) The monthly gas volume (including GEV of condensate where appropriate) received from a tied-in measured gas source (and any other receipts) must be subtracted from the total monthly battery gas volume (including GEV of condensate where appropriate) to determine the monthly battery gas production volume.

3) The table below indicates when measurement by difference may be acceptable by exception and when submission of an application may be required.

Prorated gas flow rate (excluding all measured gas source)	R* ratios		
$\leq 0.5\ 10^3\ m^3/d$	< 1.00		
$>0.5\ 10^3\ m^3/d$	≤ 0.35	> 0.35	> 0.75
Application required	**No**	**Yes**	**Yes** (Approval granted only in exceptional circumstances)

*R = Ratio of volume of all tied-in measured gas volumes (including GEV of condensate where applicable) to the total battery gas disposition volume (including fuel, flare, and vent volumes).

If any measured gas source will be tied in to an oil battery:

1) The gas and liquids from the tied-in gas source(s) must be separately and continuously measured.

2) The monthly gas volume (including GEV of condensate where appropriate) received from a tied-in measured gas source (and any other receipts) must be subtracted from the total monthly battery gas volume to determine the monthly battery gas production volume. (See Table 5.1 for options.)

3) If liquid condensate is received from a tied-in measured gas source, the monthly liquid condensate volume where appropriate must be subtracted from the total monthly oil disposition (plus/minus inventory changes and minus any other receipts) to determine the monthly battery oil production volume. (See Table 5.1 for options.)

4) The table below indicates when measurement by difference may be acceptable by exception and when submission of an application may be required.

Prorated gas flow rate (excluding all measured gas source)	R* ratios		
≤0.5 10³ m³/d	< 1.00		
>0.5 10³ m³/d	≤ 0.35	> 0.35	> 0.80
Application required	**No**	**Yes**	**Yes** (Approval granted only in exceptional circumstances)

* R = Ratio of volume of all tied-in measured gas volumes (including GEV of condensate where applicable) to the total battery gas disposition volume (including fuel, flare, and vent volumes).

Where a measured gas source will be tied into a single well battery, as shown in Figure 5.6, this situation does not qualify for an exception, and an application must be submitted to and approved by the ERCB prior to implementation.
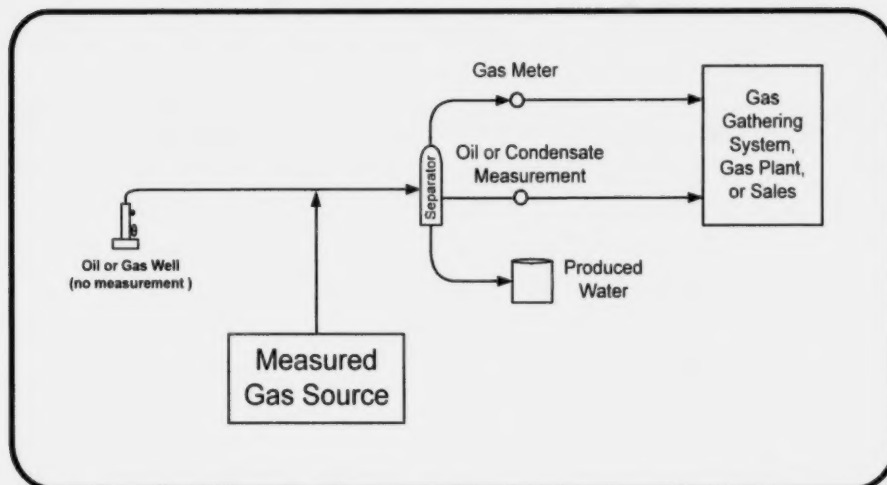
Figure 5.6

Oil Measurement by Difference

For oil streams, measurement by difference can include but is not limited to the following situations.

- Measured oil and/or oil-water emulsion from a battery delivering into an oil proration battery by truck (Figure 5.7):
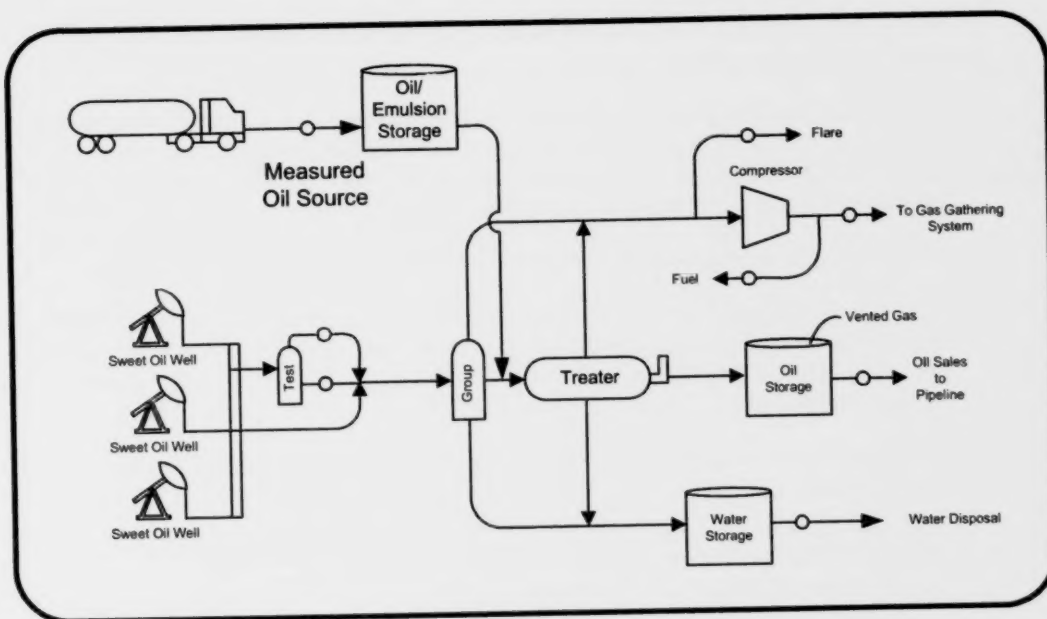
Figure 5.7

- Measured oil and/or oil-water emulsion (and gas if applicable) under pressure from a battery delivering into an oil proration battery by pipeline (Figure 5.8):
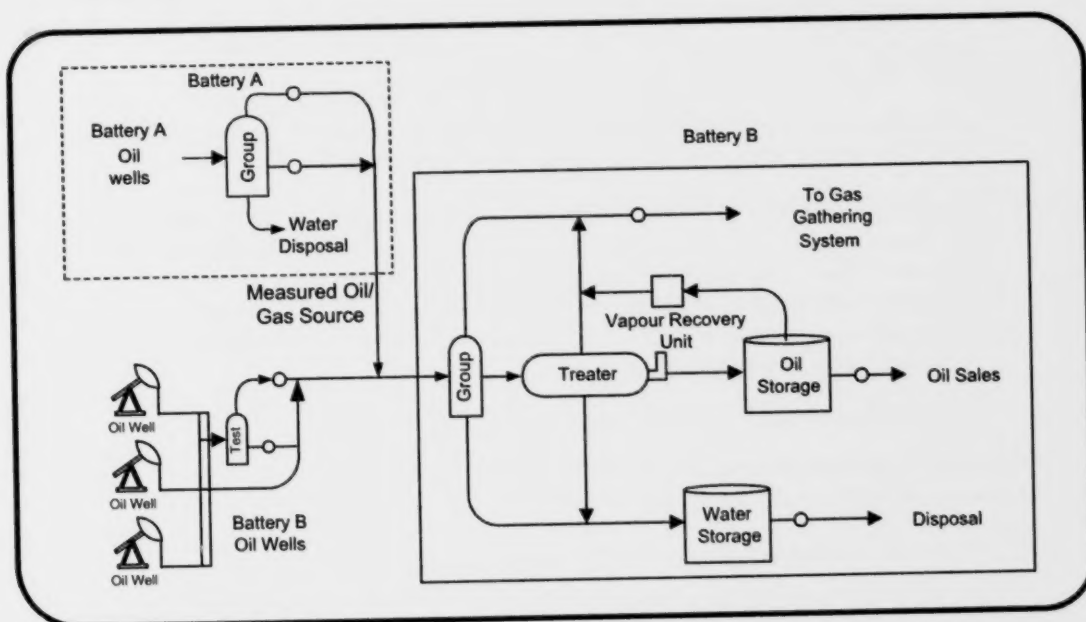
Figure 5.8

If any measured oil and/or oil-water emulsion source will be delivered to a battery including trucked-in volumes:

1) Measured oil and/or oil-water emulsion delivery/receipt volumes must be determined using equipment and/or procedures that meet delivery point measurement uncertainty requirements. In the case of oil-water emulsions, the measurement uncertainty requirements apply to total volume determination only.

2) Measured oil volumes must be determined and reported at stock tank conditions.

3) The liquids received from the delivered measured oil and/or oil-water emulsion source(s) must be subtracted from the total monthly oil and water disposition volumes (plus/minus inventory changes and minus any other receipts) to determine the monthly battery oil production volume.

4) The table below indicates when measurement by difference may be acceptable by exception and when submission of an application may be required.

Measured oil delivery/receipt volume	Application required
≤ 1000 m^3/month or if > 1000 m^3/month and ≤ 25% of monthly battery oil production volume	No
> 1000 m^3/month and > 25% of monthly battery oil production volume	Yes

5) Consideration should be given to incorporating the piped-in measured oil and/or oil-water emulsion source(s) as a satellite of the battery (if the battery is an oil proration battery) and including it in the battery's proration system. In that case, measurement by difference could be avoided.

5.5.1 Exceptions

Measurement by difference is allowed without ERCB site-specific approval if all of the applicable criteria below are met and no application is required.

If the measurement by difference will involve existing production, initial qualifying flow rates must be based on average daily flow rates (monthly flow rate divided by number of production days in the month) recorded during the six months prior to implementation of the measurement by difference. If new measured production is to be connected to a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions.

5.5.1.1 Exceptions for All Measured Gas Streams

For measured gas source(s) from either gas or oil batteries tied into a gas proration battery or an oil battery:

5.5.1.1(a) Initial Qualifying Criteria

1) Volumetric criteria for measured gas tying into a proration battery

Prorated gas flow rate (excluding all measured gas source)	R*
≤ 0.5 10^3 m^3/d	< 1.00
> 0.5 10^3 m^3/d	≤ 0.35

*R = Ratio of volume of all tied-in measured gas (including GEV of condensate where applicable) to the total gas disposition volume from the receiving battery (including fuel, flare and vent volumes).

Example:
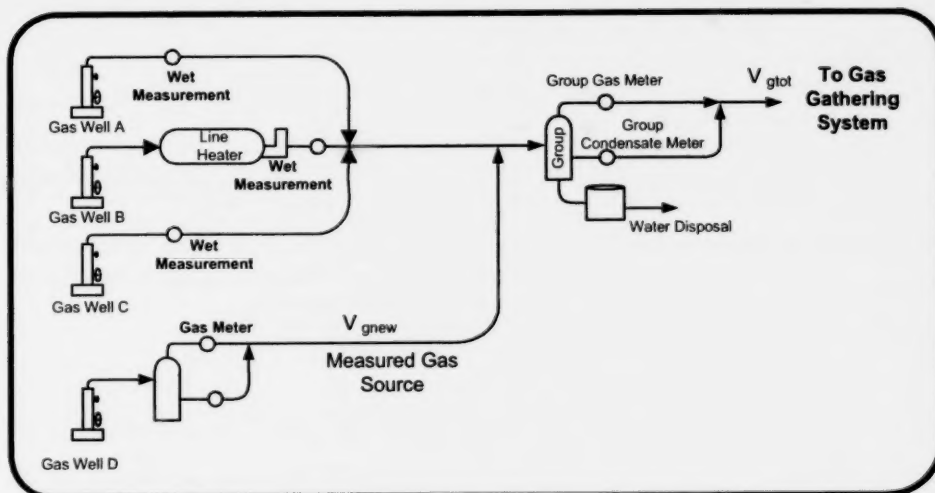
Figure 5.9

For the gas battery example in Figure 5.9,

V_{gtot} = 100 10^3 m^3/d (total of measured gas and GEV of condensate delivered out of the battery, including volumes received from Gas Well D)

V_{gnew} = 30 10^3 m^3/d (total of measured gas and GEV of condensate delivered to the battery from Gas Well D)

Prorated gas flow rate = $V_{gtot} - V_{gnew}$ = 100 – 30 = 70 10^3 m^3/d

R = 30/100 = 0.3

Since the prorated flow rate is above 0.5 10^3 m^3/d and R is below 0.35 for the Well D tie-in, it is within the acceptable exception range.

2) All proration wells flowing to the battery have common ownership and either common Crown or Freehold royalty.

   - If there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received.

   - If there is no common Crown or Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalty involved, the licensee must apply to the ERCB for approval.

3) The gas and liquid phases from the tied-in measured gas source(s) are separately and continuously measured.

4) Gas volumes received at a gas battery from the tied-in measured gas source(s) include the GEV of the measured condensate volumes if the condensate is recombined with the measured gas volumes from the new tied-in gas source.

5) If the tied-in measured gas source(s) produces condensate and it is to be connected to an oil battery, the licensee must choose from the applicable condensate delivery/reporting options in Table 5.1 below.

Table 5.1. Criteria for condensate delivery to an oil battery

Condensate rate (from measured gas sources)	≤ 2.0 m³/day and/or ≤ 5.0% of total prorated oil production	> 2.0 m³/day and/or > 5.0% of total prorated oil production
Option 1	Deliver condensate to the oil battery and report it all as liquid condensate if most of the condensate will not be flashed during the oil treatment process (minimal pressure drop and/or temperature increase).	Deliver condensate elsewhere; do not recombine it with the gas volumes delivered to the oil battery.
Option 2	Deliver condensate to the oil battery and report it all as a GEV if most of the condensate will be flashed during the oil treatment process (large pressure drop and/or temperature increase).	Tie in the gas well gas and condensate stream downstream of oil battery gas disposition measurement.
Option 3	Deliver condensate elsewhere; do not recombine it with the gas volumes delivered to the oil battery.	

6) If condensate from a tied-in measured gas source will be delivered to an oil battery and reported as a liquid volume, it is measured with a meter proved to stock tank conditions.

7) If condensate from a tied-in measured gas source will be delivered to an oil battery and reported as a GEV, it is measured with a meter proved to flow line conditions.

8) In the case of an oil battery or a gas proration battery, the monthly gas volume (including GEV of condensate where appropriate) received from a tied-in measured gas source (and any other receipts) is subtracted from the total monthly battery gas volume (including GEV of condensate where appropriate) to determine the monthly battery gas production volume.

9) In the case of an oil battery, the monthly liquid condensate volume (where appropriate) received from a tied-in measured gas source is subtracted from the total monthly oil disposition (plus inventory changes, shrinkage, if applicable, and minus any other receipts) to determine the monthly battery oil production volume.

10) If oil and/or oil-water emulsion from a tied-in measured gas source will be delivered to a gas proration battery, the total volume does not exceed 2 m³/day, it is measured with a meter proved to stock tank conditions, and it is reported as a liquid.

5.5.1.1(b)   Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) The gas and liquid phases from the tied-in measured gas source(s) were not separately and continuously measured.

2) Measurement, accounting, and reporting of condensate in items 4 to 9 under Initial Qualifying Criteria above were not followed.

3) Measurement, accounting, and reporting of oil and/or oil-water emulsion in item 10 under Initial Qualifying Criteria above were not followed.

Base measurement requirements must be reinstated if the exception is revoked due to any of the above.

5.5.1.2   Exception for Measured Oil Streams Received by Truck

5.5.1.2(a)   Initial Qualifying Criteria

1) Oil and/or oil-water emulsions may be trucked into an oil battery provided the total measured oil receipt volume is ≤ 1000 m^3/month, or if > 1000 m^3/month, the total measured oil receipt volume is $\leq 25\%$ of the monthly battery oil production volume.

2) The monthly battery oil and water production volumes are determined by subtracting the monthly measured oil and water receipt volumes from the total monthly battery oil and water disposition volumes (plus inventory change and minus any other receipts).

5.5.1.2(b)   Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) Trucked-in oil receipt was > 1000 m^3/month and the total measured oil receipt volume was $> 25\%$ of the monthly battery oil production volume.

2) The accounting methodology in item 2 under Initial Qualifying Criteria above was not followed.

There are three options to follow after revocation:

1) truck all oil receipts over 1000 m^3/month elsewhere, or

2) set up another treater train with separate inlet measurement, tankage, and outlet measurement to process the truck-in receipts prior to commingling with the battery production, or

3) obtain ERCB special approvals to continue.

5.5.1.3 Exception for Measured Oil Streams Received by Pipeline

5.5.1.3(a) Initial Qualifying Criteria

1) Oil and/or oil-water emulsions may be pipelined into an oil battery provided that the total measured oil receipt volume is < 1000 m^3/month or if > 1000 m^3/month, the total measured oil receipt volume is ≤ 25% of the monthly battery oil production volume.

2) All wells that belong to the oil battery and the tied-in oil source(s) have common ownership and either common Crown or Freehold royalty:

   • If there is no common ownership, written notification has been given to all working interest participants, with no resulting objections.

   • If there is no common Crown or Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to the ERCB for approval.

3) The monthly battery oil and water production volumes are determined by subtracting the monthly measured oil and water receipt volumes from the total monthly battery oil and water disposition volumes (plus inventory change and minus any other receipts).

4) If the measured gas from a measured live oil/emulsion production source is also commingled with the production at an oil battery, the exception criteria for gas measurement by difference are also met.

5.5.1.3(b) Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) Piped in oil receipt was > 1000 m^3 and the total measured oil receipt volume was > 25% of the monthly battery oil production volume.

2) The accounting methodology in item 3 under Initial Qualifying Criteria above was not followed.

3) Gas measurement by difference exception criteria in item 4 under Initial Qualifying Criteria above was not met.

There are three options to follow after revocation:

1) pipe all oil receipts over 1000 m^3/month elsewhere, or

2) set up another treater train with separate inlet measurement, tankage, and outlet measurement to process the pipeline receipts prior to commingling with the battery production, or

3) obtain ERCB special approvals to continue.

Oil System Example:
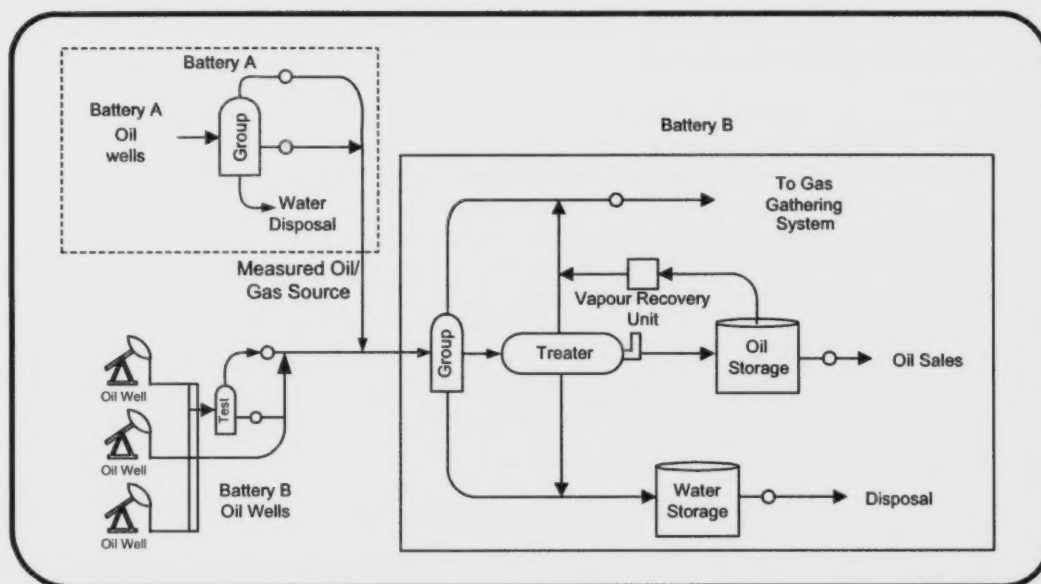
Figure 5.11

Note that with the addition of Battery A production, if the measurement by difference meets all the initial qualifying criteria and the total oil delivery volume at Battery B is over 100 m^3/d, the delivery volume must be determined by a measurement device(s) and/or procedures having $\pm 0.5\%$ uncertainty, which might require changes in measurement equipment and/or procedures at Battery B.

For this example (Figure 5.11), given the following data,

Battery A oil production volume = 20.0 m^3/d

Battery B oil production volume = 90.0 m^3/d before tying in Battery A

Battery A gas production volume = 15.0 $10^3 m^3/d$

Battery B gas production volume = 20.0 $10^3 m^3/d$ before tying in Battery A

Step 1: Calculate the monthly measured oil volume from Battery A delivered to the proration battery (Battery B) and the percentage of the prorated oil production:

Monthly measured oil production volume from Battery A = 20.0 m^3/d x 30 days = 600 m^3

Battery A oil volume as a percentage of Battery B oil production volume = 20 m^3/d / 90.0 m^3/d = 22.2%

Step 2: Calculate the R ratio for the commingled gas:

R = 15.0/(15.0 + 20.0) = 0.43

Since the Battery A monthly measured oil volume is below 1000 m^3/month, the oil volumetric criteria are met. However, the gas R ratio is over the 0.35 limit, so an application would be required.

5.5.2 Applications

The following information must be submitted with an application to add measured gas or oil/emulsion sources to a proration battery if the above criteria are not met:

1) all of the information listed in Section 5.2: Site-specific Approval Applications;

2) a discussion of the stage of depletion for pools involved, and the impact of any reduction in well measurement accuracy that may result from measurement by difference as it relates to reservoir engineering data needs; discussion of this matter by the proponent with its own reservoir engineering staff or knowledgeable external personnel is required and must be addressed in the application;

3) if there are no common Crown or Freehold royalties or common ownership, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold royalty holders and working interest participants, with no resulting objection received.

5.5.3 Considerations for Site-specific Approval

1) There are minimal equity, royalty, and reservoir engineering concerns.

2) Economic considerations: Would implementation of a proration system reduce costs enough to significantly extend operations? Have other options been considered?

3) The gas and liquids from the tied-in measured source(s) must be separately and continuously measured.

4) If the tied-in measured gas source(s) produces condensate and it is to be connected to an oil battery, the licensee must choose from the following applicable condensate delivery/reporting options:

Table 5.2. Options for condensate delivery to an oil battery

Condensate rate (from measured gas sources)	≤ 2.0 m^3/day and/or $\leq 5.0\%$ of total prorated oil production	> 2.0 m^3/day and/or $> 5.0\%$ of total prorated oil production
Option 1	Deliver condensate to the oil battery and report it all as liquid condensate if most of the condensate will not be flashed during the oil treatment process (minimal pressure drop or temperature increase).	A laboratory analysis, or a computer software simulation based on the condensate composition must be performed to determine the amount of the condensate that will be flashed during the oil treatment process. **Flashed condensate:** report it as a GEV of condensate and include it with the gas delivery/receipt volume. **Unflashed condensate:** report it as a liquid condensate delivery/receipt volume.
Option 2	Deliver condensate to the oil battery and report it all as a GEV if most of the condensate will be flashed during oil treatment process.	Deliver condensate elsewhere and do not deliver it to the oil battery.
Option 3	Deliver the condensate elsewhere and do not deliver it to the oil battery.	

5) In the case of an oil battery or a gas proration battery, the monthly gas volume (including GEV of condensate where appropriate) received from a tied-in measured gas source (and any other receipts) must be subtracted from the total monthly battery gas volume (including GEV of condensate where appropriate) to determine the monthly battery gas production volume.

6) In the case of an oil battery, the monthly liquid condensate, or oil, and/or oil-water emulsion volume (where appropriate) received from a tied-in measured source must be subtracted from the total monthly oil and/or water disposition (plus/minus inventory changes and minus any other receipts) to determine the monthly battery oil and/or water production volume.

5.6 Surface Commingling of Multiple Gas Zones/Wells

If gas wells have been completed in multiple zones and those zones are segregated in the wellbore and produced separately to surface or if there are multiple individual gas wells on the same surface location, production from each zone or each well usually has to be measured separately prior to commingling. Where applicable, such zones or wells may be commingled at surface prior to the combined production being measured, if the conditions in the "Exceptions" section below are met or upon approval of an application. Proportionate monthly production volumes must still be determined and reported for each zone/well, in accordance with the applicable procedures described below.

The following requirements do not apply to zones that qualify for the SE Alberta Shallow Gas Proration Battery procedures if specific zones are approved (without application) for commingling in the wellbore.

Commingling of zones in the wellbore require approval from the ERCB Resources Applications Group.

5.6.1 Exceptions

Surface commingling of two gas zones in a gas well or separate gas wells on the same surface location prior to measurement is allowed without ERCB site-specific approval if all the initial qualifying criteria below are met and no application is required.

Initial Qualifying Criteria

1) Both zones/wells have common ownership and common Crown or Freehold royalty:

   • If there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received.

   • If there is no common Crown or Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to the ERCB for approval.

2) Monthly average of total liquid production from both zones is $\leq 2 \ m^3/d$.

3) The combined daily flow rate of both zones/wells is $\leq 16.9 \ 10^3 \ m^3$, including GEV of condensate (if recombined). If the zones/wells to be commingled will involve existing production, initial qualifying flow rates are based on monthly average flow rates recorded during the six months prior to implementation of the commingling. If new zones/wells

are to be commingled, the initial qualifying flow rates are based on production tests conducted under the anticipated operating conditions.

4) Shut-in wellhead pressure of the lower pressure zone/well is ≥ 75% of the shut-in wellhead pressure of the higher-pressure zone.

5) The combined production from both zones/wells is measured continuously. Separation before measurement is required for both phases.

6) Check valves are installed on each flow line upstream of the commingling point.

7) Testing requirements:

- Each zone/well must be tested once per month for the first six months after commingling, then annually thereafter, and/or immediately following any significant change to the producing conditions of either zone/well.

- The tests must be conducted for at least 24 hours and must involve the separation and measurement of all gas and liquid production.

- If condensate is recombined with the gas production of the commingled zones/wells, a sample of the condensate must be taken annually and analyzed and used to determine the factor to be used to determine the GEV.

- The tests for both zones/wells must be done consecutively with stabilization periods.

- Any of the three test methods described below may be used. However, methods (i) and (ii) below are preferred, because the testing is conducted under normal flowing conditions without shutting in zones/wells, so that minimal stabilization time is required.

  i) Test taps must be installed upstream of the commingling point but downstream of the check valve so that a test separator unit can be hooked up to test each zone/well individually (Figure 5.12).

Test Method (i)
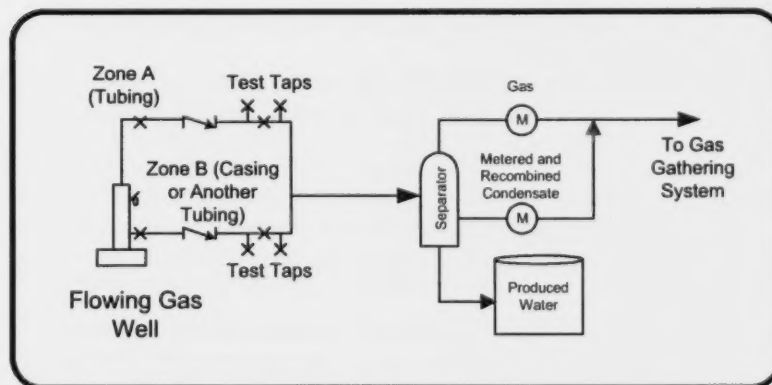
Figure 5.12

ii) Install permanent bypasses or taps to hook up temporary bypasses downstream of the check valve so that one zone/well will be bypassing the existing separation and metering equipment while the other zone/well is tested using the existing equipment. Note that the production from the bypassed zone/well must be estimated based on the production test rates (Figure 5.13).

Test Method (ii)
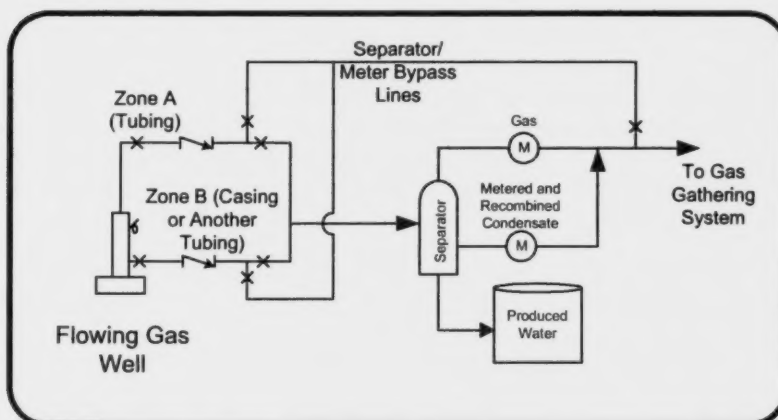
Figure 5.13

iii) Shut in one producing zone at a time and use the existing separation and measurement equipment to test each zone/well individually after stabilization.

8) The production rates determined for each zone/well by the periodic tests must be used to estimate the monthly production for each zone/well from the date they are conducted until the next test is conducted. The monthly measured combined production must be prorated to each zone/well based on the estimates, and those prorated volumes must be reported as the monthly production for each zone/well.

Revocation of Exceptions

If any of the following exists or occurs, the exception is revoked:

1) The combined production from both zones/wells was not measured continuously or there was no separation before measurement.

2) Check valves were not installed on each flow line upstream of the commingling point.

3) Testing requirements in item 7 under Initial Qualifying Criteria above were not followed.

4) The gas proration methodology in item 8 under Initial Qualifying Criteria above was not followed.

Base measurement requirements must be reinstated if the exception is revoked due to any of the above.

5.6.2 Applications

The following information must be submitted with an application to commingle production at surface prior to measurement from multiple zones in a gas well or multiple wells on the same surface location if the above criteria are not met:

1) all of the information listed in Section 5.2: Site-specific Approval Applications;

2) shut-in and proposed operating pressures at the wellhead for all zones/wells;

3) operating pressure for the gathering system at the well site measurement point;

4) proposed testing procedures to determine the individual zone/well production rates;

5) proposed accounting procedures for prorating total volumes to the individual zones/wells; and

6) if there are no common Crown or Freehold royalties or common ownership, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold royalty holders and working interest participants, with no resulting objection received.

5.6.3 Considerations for Site-specific Approval

1) Generally, there is ≤ 2 m^3/day of total liquid production from all zones/wells.

2) All zones/wells must be classified as gas zones/wells.

3) There are minimal equity, royalty, and reservoir engineering concerns.

4) The combined production of all zones/wells must be continuously measured. If there are gas and liquid components, they must be separately measured.

5) Check valves must be in place on the flow line upstream of the commingling point.

6) Testing requirements:

   - Each zone/well must be tested once per month for the first six months after commingling, then annually after that, and/or immediately following any significant change to the producing conditions of either zone/well.

   - The tests must be conducted for at least 24 hours in duration and must involve the separation and measurement of all gas and liquid production.

   - If condensate is recombined with the gas production of the commingled zones/wells, a sample of the condensate must be taken annually and analyzed and used to determine the factor that will be used to determine the GEV.

   - The tests for all zones/wells must be done consecutively, with stabilization periods.

   - Any of the three test methods described in the exceptions section above may be used, with the consideration that more than two zones/wells may be involved. However, methods (i) and (ii) are preferred, because the testing is conducted under normal flowing conditions without shutting in zones/wells, so that minimal stabilization time

is required. The ERCB may specify test procedures if specific circumstances warrant them.

7) The production rates determined for each zone/well by the periodic tests must be used to estimate the monthly production for each zone/well from the date they are conducted until the next test is conducted. The monthly measured combined production must be prorated to each zone/well based on the estimates, and those prorated volumes must be reported as the monthly production for each zone/well.

5.7   Gas Effluent Wells Exemption and Reduction for Testing Frequency

For most wells, the required minimum well testing frequency is annual unless the criteria below are met to reduce this frequency to triennial or, possibly, to complete exemption. If these criteria are not met, the licensee may still apply to the ERCB for a change to the minimum well testing frequency. The criteria below provide details of situations for which an exemption or reduction can be implemented automatically without approval from the ERCB. All other situations require approval from the ERCB.

An application for a reduction or exemption is required if the production is from **coalbed methane wells or wells producing gas from above the base of groundwater protection,** even if the initial qualifying criteria are met.

An application for reduction or exemption from the annual testing requirements is not required if the criteria listed in 5.7.1.1 or 5.7.2.1 are met. The licensee must retain the data and documentation to support the initial qualifying criteria and the last three testing records for as long as the reduction or exemption is in place. The ERCB may revoke an exemption or testing frequency reduction if an audit or inspection reveals a lack of adequate supporting data or documentation.

5.7.1   Effluent Gas Well Testing Exemption

This exemption is on a per-battery basis and applies to low-production or low-pressure pools. All wells in an effluent proration battery are exempt from the ECF-WGR testing requirements provided that the initial qualifying criteria below are met and no application is required. This exemption is revoked if any of the "Revocation of Exemption" criteria set out below apply. If any well is added to the battery, the entire battery must be re-evaluated according to the initial qualifying criteria.

The ECF for these wells are to be set at 1.00000, so that the well's estimated gas production is equal to the volume measured by the effluent meter. The well's WGR for estimating water production must be the one calculated from the last WGR test for that well. See Section 7.4 for an explanation of how to apply the ECF and the WGR.

5.7.1.1 Initial Qualifying Criteria

1) All wells flowing to the battery must have common ownership, or written notification must have been given to all working interest participants with no resulting objections.

2) All wells flowing to the battery must have common Crown or Freehold royalties. If there is no common Crown or Freehold royalty and only Freehold royalties are involved, written notification must have been given to all Freehold royalty holders with no resulting objection received. If there is a mixture of Crown and Freehold royalty involved, the licensee must apply to the ERCB for approval.

3) Each well must be gas wells subject to effluent measurement and have produced for at least 30 months over the previous three years.

4) Each well must have had at least one ECF-WGR test in each of the previous three years.

5) The total liquid to gas ratio (LGR) for each of the ECF-WGR tests during the previous three years must have been less than 0.05 m^3 liquid / 10^3 m^3 gas. LGR is calculated by dividing the total water and condensate test volumes by the measured test gas volume.

6) Excluding all shut-in periods from the calculations, one of the following must apply:

   • no well has a monthly average actual gas production rate (including gas equivalent of condensate) for the last 12 months that exceeds 6.0 10^3 m^3/d, or

   • the maximum normal operating pressure at the effluent meter for all wells has not exceeded 280 kilopascals gauge (kPag) for any month out of the last 12 months (this does not apply to pressure increases due to compressor shutdown or other operational upsets).

5.7.1.2 Revocation of Exemption

The exemption is revoked and annual ECF-WGR testing of all wells must recommence if any of the following occurs:

1) A re-evaluation of the entire battery after a well(s) is added does not meet the initial qualifying criteria.

2) The working interest participants for any well flowing to the battery have changed and a new working interest participant objects to the exemption.

3) Any well within the battery has exceeded the 6.0 10^3 m^3/d monthly average actual gas production rate (including gas equivalent of condensate), or the maximum normal operating pressure at the effluent meter for any well in the battery exceeds 280 kPag.

4) An artificial lift system is put in place for any well in the battery. Examples include plunger lifts, on/off controls, pumpjacks, and gas lifts.

5.7.2   Testing Frequency Reduction for Effluent Gas Wells Producing Low Rates of Water

The ECF-WGR test frequency for wells in an effluent proration battery may be reduced to a minimum of once every three years on a per-well basis provided that the initial qualifying criteria below are met and no application is required. This reduction is revoked if any of the "Revocation of Frequency Reduction" criteria set out below apply.

5.7.2.1 Initial Qualifying Criteria

1) All wells flowing to the battery must have common ownership, or written notification must have been given to all working interest participants with no resulting objections.

2) All wells flowing to the battery must have common Crown or Freehold royalties. If there is no common Crown or Freehold royalty and only Freehold royalties are involved, written notification must have been given to all Freehold royalty holders with no resulting objection received. If there is a mixture of Crown and Freehold royalty involved, the licensee must apply to the ERCB for approval.

3) Each well must be gas wells subject to effluent measurement and have produced for at least 30 months over the previous three years.

4) Each well must have had at least one ECF-WGR test in each of the previous three years.

5) For the ECF-WGR tests for each of the wells that qualify for this reduction during the previous three years, one of the following must apply:

- the LGR is less than 0.05 m^3 liquid /10^3 m^3 gas, or

- the LGR is greater than 0.05 m^3 liquid/10^3 m^3 gas, the water content does not exceed 0.05 m^3 water /10^3 m^3 gas at separator conditions, and all LGR test data points for the previous three years are within ± 0.0% of the average of those test data points. If the well qualifies for this reduction, the three most recent data points are to be used to calculate the average on a go-forward basis, and all data points must be within ± 10.0% of that average (see example below).

Example:

	Sept. 00	Sept. 01	Mar. 02	Sept. 02	Mar. 03	3-yr. average	Max. dev.
Well LGR	0.120	0.128	0.117	0.110	0.131	0.1212	
Dev. from average	1.0%	5.6%	3.5%	9.2%	8.1%		9.2%

Since the maximum deviation is within 10% of the three-year average, this well qualifies for reduction to testing once every three years.

When another test is conducted in 2006, the average is calculated as below:

	Sept. 02	Mar. 03	Mar. 06	3-data-point average	Max. dev.
Well LGR	0.110	0.131	0.120	0.1203	
Dev. from average	8.6%	8.9%	0.3%		8.9%

Since the maximum deviation is within 10% of the last three-data-point average, this well continues to qualify for reduction.

5.7.2.2 Revocation of Frequency Reduction

The frequency reduction is revoked for the individual well(s) and annual ECF-WGR testing must recommence if any of the following occurs:

1) The working interest participants for any well flowing to the battery have changed and a new working interest participant objects to the reduction.

2) Wells under frequency reduction were not tested as required.

3) The liquid water content of a well at separator conditions has exceeded the 0.05 m^3 water/ 10^3 m^3 gas rate.

4) The LGR exceeds 0.05 m^3 liquid / 10^3 m^3 gas and the LGR is not within ± 10.0% of the average of the three most recent test data points.

The licensee should monitor the gas and water production for changes such as the following:

- changes in water proration factors for the effluent battery beyond the normal operating range (see Section 3.1 of this directive)

- changes to spiking chart traces or electronic flow measurement at the meters
- increases in intensity of liquid slugging

5.7.3   Applications

The following information must be submitted with an application for exemption or reduction in testing frequency for an effluent gas well(s) if the above criteria are not met:

1) all of the information listed in Section 5.2: Site-specific Approval Applications;

2) a list of all the wells, including from which formations, where commingled production has occurred in the wellbore;

3) a list of all wells and corresponding pool(s) broken down into the following groups:
    - in the ERCB-designated oil sands areas (include a sublist of the wells and the corresponding pool[s] associated with bitumen),
    - coalbed methane wells, and
    - producing from above the base of groundwater protection;

Optional information to be submitted upon request by the ERCB

4) map(s) of the producing pool(s) in item 2, showing
    a) all the wells that have produced from them but distinguishing between the application and non-application wells, and
    b) any area in the pool(s) where underlying water exists;

5) a discussion of the testing method currently used and, if different, the method proposed to be used to determine water production;

6) a discussion on the initial location of the gas/water interface, including any movement in it during the past and where it is likely to reside in the predicted future; explain the potential impact on water production due to the position and movement of the gas/water interface;

7) a discussion on the susceptibility of the producing formation to water damage, the impact on ultimate recovery, and how historical and future operations will address the potential for conservation losses resulting from water production problems;

8) a discussion on the source of produced water and how this was determined;

9) a list of the wells and corresponding pool(s) broken down according to where
    a) water coning could explain increased water production,
    b) shut-in or suspension has occurred at any time for more than six months (with the reasons why),
    c) abandonment has occurred (with the reasons why), and
    d) production has not exceeded three years;

10) a historical production plot for any well in items 2 and 3 that exceeds a WGR of $0.05 \ m^3/10^3 \ m^3$ at any time during its producing life:

    a) this plot should be on an individual well basis with time on the x-axis and water rate, gas rate, and WGR plotted together on the y-axis,

    b) the units for time should be in years, gas rate in $10^3 \ m^3/d$, water rate in m^3/d, and WGR in $m^3/10^3 \ m^3$, and

    c) the plot should show the complete producing life of the well, and the production should be based on data submitted to the ERCB using calendar time;

11) a discussion on where the water may be coming from for the wells plotted in item 10, and an explanation of any significant changes in water rate or WGR data shown on these plots; and

12) an estimate of the remaining recoverable gas reserves for the pool(s) and the average gas production rate over the last year for each producing well in the application.

6 Conventional Oil Measurement

This section presents the base requirements and exceptions for conventional crude oil and emulsion measurements from wells and batteries in the upstream oil and gas industry that are used in determining volumes for reporting to the PRA. The requirements for crude oil/emulsion volumes transported by truck are detailed in *IL 90-6* and *IL 92-8*, which will be replaced by the Trucked Liquid Measurement section (under development).

Conventional crude oil has the following characteristics:

- it is a mixture mainly of pentanes and heavier hydrocarbons that may be contaminated with sulphur compounds,
- it is recovered or is recoverable at a well from an underground reservoir,
- it is liquid at the conditions under which its volume is measured or estimated, and
- it must have a density of less than 920 kg/m^3 at standard conditions.

6.1 General Requirements

Crude oil may be found in association with water in an emulsion. In such cases, the total liquid volume must be measured, and the relative volumes of oil and water in the emulsion must be determined by obtaining and analyzing a representative sample of the emulsion, by using a product analyzer, or by other means if applicable. Applications for which estimation of water content is appropriate (e.g., inventory) are covered in more detail later in this section.

A licensee must measure produced crude oil/emulsion volumes by tank gauging, weigh scale, or meter, unless otherwise stated in this directive. The ERCB will consider an oil measurement system to be in compliance if the base requirements detailed below are met. The ERCB may stipulate additional or alternative requirements for any specific situation based on a site-specific assessment and will inform licensees in writing of any additional or alternative requirements respecting their facilities.

6.2 General Measurement, Accounting, and Reporting Requirements for Various Battery Types

6.2.1 General Accounting Formula

Production = Total disposition + Closing inventory – Opening inventory – Total receipts

6.2.2 Oil Batteries

6.2.2.1 General

All wells in the battery must be classified as oil wells.

Liquid production from an oil battery must be measured as an oil, water, or oil/water emulsion volume. This measurement may be performed at the battery site, a truck delivery/receipt point, or a pipeline delivery point. The meter factor obtained from meter proving must be applied to the meter volumes until another prove is conducted.

All wells in a multiwell oil battery must be subject to the same type of measurement (measured or prorated). If there is a mixture of measured and prorated wells within the same battery, the exception criteria in Section 5.5 must be met or an ERCB site-specific approval must be obtained.

Production from gas batteries or other oil/bitumen batteries may not be connected to an oil proration battery upstream of the oil proration battery group measurement point(s) unless specific criteria are met or an ERCB approval is obtained (see Section 5.5). For oil delivered to a gas system, see Section 6.2.3.1.

Any oil well that produces fluids from any formation is considered on production and a battery code is required to report the production on the PRA even for a "test." See *On-line Supplement to Directive 007: Volumetric and Infrastructure Requirements,* Appendix 8, for load fluid reporting.

6.2.2.2 Single-well Battery

Oil/emulsion must be separated from gas and measured.

6.2.2.3 Multiwell Group Battery

Each well must have its own separation and measurement equipment, similar to a single-well battery.

All equipment for the wells in the battery must share a common surface location.

6.2.2.4 Proration Battery

All well production is commingled prior to the total battery oil/emulsion being separated from the gas and measured. Individual monthly well oil production is estimated based on periodic well tests and corrected to the actual monthly volume through the use of a proration factor.

Double proration, whereby the proration oil battery disposition volume(s) is prorated to group measurement points and then further prorated to the wells, is not allowed unless an approval to do so has been obtained from the ERCB.

6.2.3 Gas Batteries Producing Oil

6.2.3.1 General

All wells in the battery must be classified as gas wells.

Oil production, receipt, disposition, and inventory volumes must be reported as liquid oil. Oil volumes must not be converted to a gas equivalent volume (GEV) and must not be added to the gas volumes. If oil is recombined with the gas and delivered to a gas plant (through a gas gathering system), the oil volume as determined at the battery must be reported as delivered to the gas plant. The gas plant must report the oil disposition as appropriate.

6.2.3.2 Single-well Battery

Oil/emulsion must be separated from gas and measured.

6.2.3.3 Multiwell Group Battery

Each well must have its own separation and measurement equipment, similar to a single-well battery, and its gas production must be connected by pipeline to a common location for further processing.

6.3   Base Requirements for Oil Measurement

6.3.1   System Design and Installation of Measurement Devices

The system design and installation of oil/emulsion measurement devices must be in accordance with the following.

6.3.1.1 Meters

The meter and its associated peripheral equipment and piping must be designed and installed according to applicable industry-accepted standards or the manufacturer's recommendation. Temperature measurement must be in accordance with Table 6.1.

Table 6.1. Required frequency for oil/emulsion temperature measurement

Criteria or events	Minimum frequency
Delivery point (if a meter is used)	Continuous
EFM devices (optional for production proration testing)	
Delivery point batch volumes into a pipeline or receipt at a battery/facility using tank gauging	One reading per load

Note that the temperature-measuring element must be installed in the flow stream and be representative of the oil/emulsion stream temperature—that is, the licensee cannot use the surface temperature of the piping or install the element where there is normally no flow. A meter equipped with a temperature compensation device is considered to have continuous temperature measurement.

6.3.1.2 Tank Gauging

The licensee must ensure that the gauge tape, the strapping table, or the calculation used to convert gauge level to a liquid volume is appropriate for the tank being gauged.

With the exception of inventory determination, the licensee must ensure that the tank level is not changing when the gauge readings are taken. (This often requires isolating the tank before gauging.)

Gauge boards must not be used for delivery point measurement.

6.3.1.2(a)   Tank Sizing Requirements for Testing or Delivery Point Measurement

The relative accuracy of the volume determination using tank gauging depends on several factors and includes both random and bias errors. Such factors include the accuracy of the gauge tape, the strapping table used, the condition of the gauge point on the tank, the difference in temperature of the of the tank volume before and after the transaction, and the level of operator training.

The primary function is to convert a level measurement to a volume measurement, and the precision that can be achieved when tank gauging is of concern. Since the precision that can be achieved by manual tank gauging is relatively consistent, the main way to determine accurate tank volumes is by ensuring that the level change corresponding to the transaction or test volume is large compared to the overall tank height. This can be achieved by selecting a tank design such that the ratio of volume per height of fluid is low.

The following formulas may be used to determine these values:

$$V \geq a \times d^2 \qquad \text{or} \qquad d \leq (V/a)^{1/2}$$

where

V = test fluid volume or delivery point batch volume in m^3
d = tank diameter in metres
a = accuracy coefficient

The accuracy coefficient = 0.39 for all test fluid volumes
= 0.39 for delivery point batch volumes $\leq$ 100 m^3/d
= 0.92 for delivery point batch volumes > 100 m^3/d

6.3.1.2(b) Flow Line Delivery Point Measurement Gauging

When tank gauging is used to determine an oil/emulsion volume, the gauging procedures must be conducted in accordance with the following:

- The licensee must ensure that the gauge/strapping table used to convert the gauge level to a liquid volume is specific for the tank being gauged.

- With the exception of inventory determination, the licensee must ensure that the tank level is not changing or is stabilized when the gauge readings are taken. This often requires isolating or shutting in the tank before gauging.

- All gauge tapes and electronic level devices must have a minimum resolution of 3 mm.

Manual Tank Gauging

For tanks with a nominal capacity greater than 160 m^3, two consecutive readings within 10 mm of each other are required; the two readings are then averaged.

For tanks with a nominal capacity of 160 m^3 or less, one reading is acceptable.

Automatic Tank Gauging

One reading of the instrument is acceptable.

6.3.1.2(c) Inventory Tank Gauging

For monthly inventory measurement gauging, one reading of the gauge tape, gauge board, or automatic tank gauge is acceptable. The required resolution for reporting is to the nearest 75 mm.

Gauge board markings must be no farther apart than 150 mm.

6.3.1.2(d) Test Tank Gauging

For gauge measurement on test tanks, one reading of the gauge tape, gauge board, or automatic tank gauge is acceptable.

Gauge board markings must be no farther apart than 25 mm, and where safe work conditions allow, gauge board readings should be taken at eye level. The required resolution for reporting is to the nearest 10 mm.

6.3.1.3 Electronic Flow Measurement

Electronic oil measurement systems must be designed and installed according to the requirements in Section 6.8. Any EFM system designed and installed in accordance with API MPMS, Chapter 21.2, is considered to have met the audit trail and reporting requirements, but a performance evaluation is still required in accordance with Section 6.8.

6.3.2 Volumetric Calculations

Crude oil volume measurements must be determined to two decimal places and rounded to one decimal place for monthly reporting. Where there is more than one volume determination within the month at a reporting point, the volumes determined to two decimal places must be totalled prior to the total being rounded to one decimal place for reporting purposes.

6.3.2.1 Temperature Correction Requirements

Temperature measurement used for volume correction must be representative of the actual fluid temperature. Total monthly oil volumes for wells (production) and batteries (production, receipts, dispositions, and delivery point) must be reported in cubic metres at a standard temperature of 15°C and rounded to the nearest tenth of a cubic metre (0.1 m³). Battery or facility opening and closing inventory volumes for monthly reporting must be rounded to the nearest 0.1 m³ but do not require correction to 15°C. The temperature correction (Correction for the effect of Temperature on Liquids [CTL]) factor must be determined in accordance with API MPMS, Chapter 11.1.

In a proration battery, if well test oil volumes are determined by a meter, temperature compensation must be applied using one of the following methods:

- Apply a composite meter factor that incorporates a CTL factor. To arrive at a composite meter factor, divide the temperature corrected prover volume by the indicated meter volume for each prover run.

- Apply a CTL factor in real time using an electronic flow measurement system.

- Apply a CTL factor to the total test volume based on a single temperature measurement taken during the test.

See Table 6.1 for temperature measurement frequency requirement.

6.3.2.2 Pressure Correction Requirements

Correction to a 0.0 kPa gauge (atmospheric pressure) must be performed for continuous flow crude oil pipeline measurement where custody transfer measurement is performed. The pressure correction (Correction for the effect of Pressure on Liquids [CPL]) factor must be determined in accordance with API MPMS, Chapter 11.

All other crude oil/emulsion measurements, such as delivery point and test oil, do not require CPL correction.

6.3.2.3 Shrinkage Factor

At the test separator, the oil is normally at its bubble point (equilibrium vapour pressure) condition. When the oil is discharged to a stock tank at atmospheric condition, the light components in the oil evaporate, causing a reduction in liquid volume. Therefore, a shrinkage factor must be applied to correct the measured liquid volume from metering condition to stock tank condition. Degassing a prover prior to taking the volume reading for each run effectively incorporates the shrinkage factor into the meter factor and requires no further correction for shrinkage. However, sufficient time must be allowed for the degassing to take place.

Shrinkage factors can also be determined by laboratory or software simulation methods and applied to the test oil volume, eliminating the need for degassing during proving. Shrinkage

factors may be determined at either a well or battery level, depending on variation. The frequency of shrinkage factor determination should reflect changes in reservoir or operating conditions. Further details on calculation and use of shrinkage factors are in API MPMS, Chapter 20.1.

6.3.2.4 General Volume Calculations

The oil volume calculations must adhere to the following:

1) Total indicated volume for the transaction period (daily, weekly, monthly) is measured and recorded. This applies to measurement by meter, by weight scales, or by tank gauging.

2) The volumetric meter factor for the flow meter is applied to the total indicated volume.

3) The percentage of water in the gross volume is determined by measuring the percentage of sediments and water (% S&W) of a representative sample or by continuous on-line measurement. The result is a quantified volume of oil and of water. It is also possible to use a computer algorithm to determine the oil and water volumes in the emulsion based on the measured density of the emulsion and the known density of the oil and water components of the emulsion. The oil and water base density must be based on an analysis of the actual oil and water production being measured and must be corrected for the temperature at which the emulsion density is measured, and these densities must be updated annually or more frequently if there is a change in reservoir conditions of the well.

4) A shrinkage factor is applied to the oil volume in order to determine the volume at stock tank conditions (ambient pressure). As noted in Section 6.3.2.3 above, some applications may already have the shrinkage factor incorporated into the meter factor.

5) For delivery point measurement, a CTL is applied to the oil volume to determine the volume at a standard temperature of 15°C. Some applications may already have the CTL incorporated into the meter factor. Applying CTL is optional for a test oil volume, as long as the selected practice is consistent for the area.

The general formula for determining oil volumes is as follows:

Oil volume = (End reading – Start reading) x MF x (1 – %S&W) x CTL x CPL x SF

where
MF	= meter factor
%S&W	= percentage of sediments and water
SF	= shrinkage factor due to apparent loss of volume due to entrained gas
CTL	= volume correction factor for the effects of temperature on liquid
CPL	= volume correction factor for the effects of pressure on liquid

For test oil measurement, composite meter factors may include CTL, CPL, or shrinkage factor or some combination of the three, depending on the application.

A composite meter factor including CPL is acceptable where pressure is stable between proves.

Correction for shrinkage is not required when oil has been reduced to atmospheric pressure prior to measurement.

6.3.3 Production Data Verification and Audit Trail

The field data, records, and any calculations or estimations, including EFM, relating to ERCB-required production data submitted to the PRA must be kept for inspection upon request. The reported data verification and audit trails must be in accordance with the following:

1) Test records: any records and documentation produced in the production proration testing of wells that affect measured volumes

2) Proving records: any records and documentation produced in the proving of meters and calibration of the prover and all peripheral devices (if the prover and peripheral devices are owned and operated by the licensee)

3) S&W records: any records and documentation produced in the determination of relative oil/water percentages that affect volumes

4) Delivery and receipt records: any records and documentation produced in the determination of delivery or receipt volumes

5) Estimation records: any records and documentation related to the estimation of reported volumes, including estimation methodology, record of event, and approvals

6) Tank gauging records: any records and documentation produced in the determination of reported volumes

7) Volume loss records: any records and documentation for volumes lost due to incidents such as theft, spills, and fires

8) EFM: any records and documentation (electronic, magnetic, or paper form) produced in the determination of measured volumes in accordance with the EFM requirements in Section 6.8

6.4 Field Operations

6.4.1 Production Hours

Physical well shut-ins and emergency shutdowns (ESDs) are considered downtime. Other occurrences resulting in downtime may be wax or hydrates plugging lines and some other failures. If the well has no oil production but still has gas production, it is considered to be on production. The operations personnel have to make a determination based on the operating environment in other situations when the wells are not physically shut in but may not have oil and gas production.

Oil wells operating on an on/off cycle basis, such as intermittent timers, pump-off controls, and plunger lifts, that are "operating normally and as designed on repeated cycles" and where part of the operation involves shutdown of pump equipment and/or periodic shut-in of the wells (as part of the repeated cycle) are considered on production even when the wells are not pumping or flowing.

6.4.2 Fluid Sampling Requirements for S&W Determination (and Density)

S&W determination procedure including the frequency of sampling must be representative of the entire volume transaction as well as the subsequent S&W sample analysis. There are two

methods to obtain this measurement: sampling or on-line analysis using a suitable instrument (water-cut analyzer or product analyzer). Sampling can be categorized by two methods: spot or grab sampling or continuous proportional sampling. It is important that the sample location be carefully selected such that the flowing stream is adequately mixed. This can be achieved by

- installing in-line mixers,
- selecting a sampling point that offers the most practical location for collecting a sample that is mixed, such as after valves, elbows, and reducers,
- selecting a sampling point that is downstream of a metering point because of the piping elements associated with a meter run, or
- collecting samples from a number of different locations, analyzing them, and making a selection based on the location that provides the most consistent and reasonable analysis.

Grab or spot sampling may be used if the water cut is below 10% for proration oil testing. Otherwise continuous proportional sampling or the use of a product analyzer is required.

Water-cut analyzers operate on a number of different principles and often are suited for specific applications. Analyzers must be installed and maintained in accordance with the manufacturer's recommendations.

For a single-well oil battery or a multiwell group oil battery delivering emulsion off-site, the volumes will be determined by the receiving facilities.

For single-well oil batteries with two-phase or three-phase separators delivering produced oil/emulsions by pipeline to another battery, the sample must be taken at or near the oil/emulsion meter using a continuous proportional sampler. Or a product analyzer is used on-site. This is a measurement-by-difference situation (see Section 5.5).

For an oil battery with emulsion tanks, the oil and water inventory volumes in the emulsion tanks may be determined by:

- taking a spot (grab) sample anywhere between the wellhead or separator and the tank and applying the %S&W to the tank inventory,
- using water-indicating paste on the gauge tape to determine the water/oil interface in the tank inventory,
- using a representative thief sample taken from the tank,
- taking the average %S&W of the total battery production and applying that to the tank inventory,
- using the average %S&W of the trucked out volumes, or
- deeming the tank inventory to be entirely oil and making changes/amendments based on delivery volumes.

6.4.3   S&W Analysis

The licensee must select the most appropriate method for determining the %S&W. There are three methods generally considered acceptable by the ERCB based on the %S&W:

1) the centrifuge or Karl Fischer method (combined with separate method for sediment determination) for water cuts between 0 and 10%,

2) the graduated cylinder method of a larger sample for water cuts between 10 and 80% and centrifuging the oil emulsion portion, and

3) the graduated cylinder method of a larger sample for water cuts between 80 and 100% and not centrifuging the oil emulsion portion.

Recommended procedures for these three methods are shown in Appendix 4. Any alternative methods must be supported by testing that shows representative results are achieved and these alternative procedures must be made available to the ERCB upon request.

6.4.4 Proration Well Testing

Proration testing requirements for conventional crude oil wells are detailed in Table 6.2.

Table 6.2. Proration testing requirements for conventional crude oil wells

Class[a]				Minimum time	
No.	Name	Oil rate (m³/d)	Minimum test frequency	between tests[b] (days)	Minimum test duration[c] (hours)
1	High	> 30	3 per month[d]	5	12
2	Medium	> 6 but ≤ 30	2 per month[e]	10	22
3	Low	> 2 but ≤ 6	1 per month	15	22
4	Stripper	≤ 2	1 every quarter	45	22

[a] Classification for each well must be determined at least semiannually based on the average daily oil rate since the last assessment. If a well experiences operational changes that cause a change in the oil rate that could affect the classification, the operator must immediately change the classification. The average daily oil rate must be based on producing days (not calendar days).

[b] Minimum separation time between tests if minimum number of tests are conducted—the time between tests may be reduced if more than the minimum number of tests are conducted.

[c] Licensees should conduct longer duration tests for wells exhibiting erratic rates to obtain more representative test data.

[d] For Class 1 wells, the minimum test frequency is based on the assumption that the well is on production for the entire calendar month. The test frequency may be reduced to two per month if the well is shut in for at least 10 days within the month and to one per month if the well is shut in for at least 20 days within the month.

[e] For Class 2 wells, the minimum test frequency is based on the assumption that the well is on production for the entire calendar month. The test frequency may be reduced to one per month if the well is shut in for at least 15 days within the month.

6.4.4.1 Well Test Considerations

If there is a change in operating conditions during a test, such as due to a power failure or a change in choke setting, the test must be rejected and a new test must be conducted.

If there is insufficient or lost test data, such as due to meter failure, the test must be rejected and a new test must be conducted.

If there is a significant change in oil, gas, or water for a test, the validity of the test should be questioned and a retest should be considered.

Sufficient purge time must be allowed to ensure that liquids from the previous test are displaced by the new test well liquids.

The pressure difference between the test separator and the group line must not exceed 200 kPa.

A well test may be stopped early for operational reasons and still be considered valid. Reasons for the short test must be documented and made available to the ERCB upon request.

6.4.4.2 Common Flow Lines

For common flow lines, a well test must be conducted, with all other wells on the common flow line shut in following adequate purge time.

Combined (cascade) testing is allowed for common flow-lined wells, provided that the conditions in Section 6.7 are met. However, the combined test must be conducted first, and then the low gas producing well must be shut in to test the high gas producing well, allowing sufficient purging and stabilization time.

6.4.4.3 Field Header and Common Flow Line Purging

If a field header is located in the same building as the test separator, the test separator must be purged by allowing at least two liquid dumps to occur prior to starting the well test. The field header must clearly identify which well is tied to the header valves.

If a field header is not located in the same building as the test separator, sufficient purge time must be allowed to ensure that liquids from the previous test are replaced by the new test well liquids.

If two or more wells are tied into a common flow line, only one well must be produced during the well test, and the other well(s) must be shut in. Similar to a field header situation, sufficient purge time must be allowed to ensure that liquids from the previous production condition are replaced by the new test well liquids.

Sufficient purge time must be calculated as follows:

Purge time = Test line volume ÷ New test well liquid flow rate

Example: Calculate the minimum purge time required for the following test line:

Test line dimensions = 1500 m length, 88.9 mm OD pipe, 3.2 mm wall thickness

Previous well test flow rates = 5.5 m^3 oil/d, 12.0 m^3 water/d

Step 1

$$d = (88.9 - 3.2 \times 2) \div 1000 = 0.0825 \text{ m}$$

$$
\begin{aligned}
\text{Test line volume} &= (3.142 \times d^2 \times \text{length}) \div 4 \\
&= (3.142 \times (0.0825)^2 \times 1500) \div 4 \\
&= 8.02 \text{ m}^3
\end{aligned}
$$

Step 2

Purge time required = Test line volume (m^3) ÷ Well flow rate (m^3/h)

$$
\begin{aligned}
\text{Well total liquid flow rate} &= (5.5 \text{ m}^3 + 12.0 \text{ m}^3) \div 24 \text{ h} \\
&= 0.729 \text{ m}^3/\text{h}
\end{aligned}
$$

$$
\begin{aligned}
\text{Purge time required} &= 8.02 \text{ m}^3 \div 0.729 \text{ m}^3/\text{h} \\
&= 11.0 \text{ h}
\end{aligned}
$$

Therefore, the minimum purge time required is 11.0 hours.

6.5 Oil Proration Battery Accounting and Reporting Requirements

Prorated production is an accounting system or procedure in which the total battery production is allocated to wells based on individual well tests. Production from multiple oil wells may be commingled before separation and continuous single-phase measurement of the components (see Figure 6.1). Individual well production must be tested periodically to determine the production rates that can be used to estimate the well's monthly production volume. The estimated monthly well production volume is corrected using a proration factor. In summary, the following must be performed (see Section 6.5.1 for details):

- test production volumes of gas (in 10^3 m^3) and oil and water (in m^3) rounded to two decimal places;
- record test duration hours to two decimal places with the nearest quarter hour as the minimum resolution;
- determine the hour production rate for each product from the well;
- determine the estimated well production by multiplying the hour rate by the monthly hours of production; and
- determine the actual (prorated) production volume by multiplying the estimated well production by the proration factor (the total actual battery production volume divided by the total estimated battery production volume).
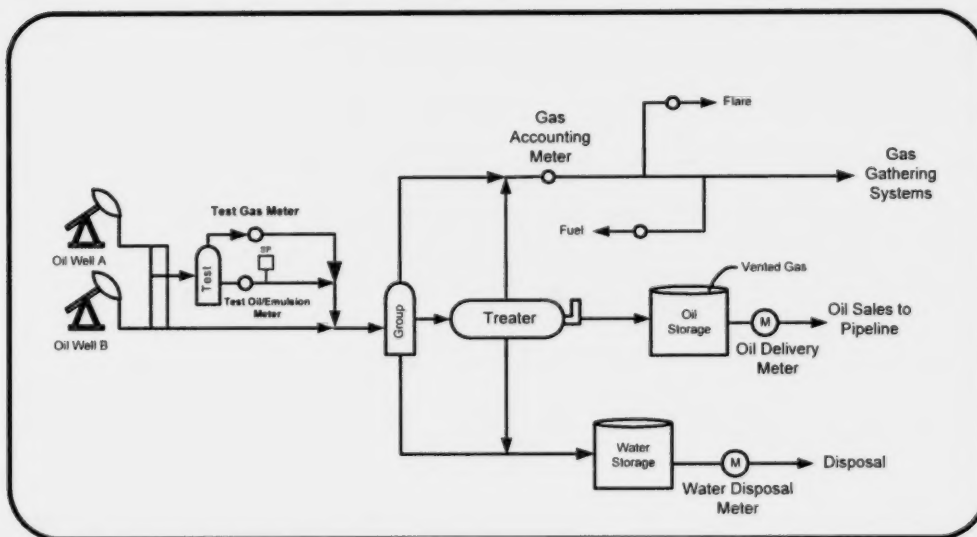
Figure 6.1. Proration testing battery

The minimum test frequency and duration requirements (see Table 6.2) apply to all conventional oil wells under primary production and waterflood operations included in proration batteries.

Monitoring the performance of miscible floods and other enhanced oil recovery schemes usually requires testing criteria other than rate alone; therefore, testing requirements for miscible flood schemes are set out in each scheme approval.

Licensees must monitor the classification for wells producing to a battery and meet the required testing frequency and duration for each well (see Table 6.2). Some latitude to the classifications is allowed. For example, considering all wells producing to a battery or

satellite as a lower rate class would be considered appropriate if a small proportion of the wells only marginally exceeded the upper limit of that classification.

Many low-rate and stripper wells exhibit erratic production rates due to high water-oil ratios or gas-oil ratios, and oversized production lines and test separators can make accurate measurement difficult. Longer test duration can improve test accuracy for many of these wells. To allow licensees the opportunity to conduct longer duration tests, class 3 and 4 wells are allowed to use up to an eight-day cycle chart drive for measurement of test gas production volumes.

The use of automatic well testing equipment and procedures with EFM provides licensees the opportunity to conduct tests of shorter durations than specified in Table 6.2. The automation computer can monitor the test and use statistical calculation methods to ensure that a representative rate is obtained prior to terminating the test. This practice is acceptable when

- the accumulated oil test volume is polled at a frequency of at least once per hour,
- the criteria for stabilization ensures that the uncertainty for the monthly well oil volume does not exceed half of the maximum uncertainty of monthly volume stipulated in Section 1, and
- the computer program is properly documented and available to the ERCB upon request.

The test-to-test method, whereby data from a test are used to estimate production until the next test is conducted, must be used to estimate the production volume from each oil well based on the test rate and the total production hours. This production estimation method and the proration methodology are outlined below. A licensee may use its own worksheet format, provided that the required data are retained and available to the ERCB upon request.

6.5.1 Proration Estimated Volume Calculation

Calculate the estimated production of each well from the test data using the sample worksheet on the next page.

1) Calculate the test rate/hour for crude oil, gas, and water:

   Rate per hour = Test production volume (including GIS volumes for gas) ÷ Test duration (hr)

   Enter the test rate/hour rounded to four decimal places.

2) Calculate the hours of production for each test rate during the reporting month. Include only the hours of prorated production:

   a) hours of production from the first day of the month to the start of the first test for the month—data from the last test conducted during the previous month will be used to estimate production until the first test for the month is conducted, and

   b) hours of production from the start of each test conducted during the month up to the start of the next test, or the end of the month, whichever is applicable.

   Enter the hours produced rounded to the nearest hour.

3) Calculate the estimated production of oil, gas, and water for the production hours applicable to each test rate:

   Estimated production = Test rate/hour x Hours produced

Enter the estimated production of oil, gas, and water rounded to one decimal place.

4) Calculate the totals for each well:

Add the hours produced that are applicable to each test rate and enter the total.

Add the estimated production of oil, gas, and water, and enter the totals.

Note that if a GOR is used to estimate the well gas production in accordance with Section 4.3.5:

Estimated well gas production = Estimated well oil production x GOR

UID	AB WI 100060100101W400						Hourly test rate				Estimated production		
	Test date		Test oil	Test gas	Test water	Test duration[c]	Oil	Gas	Water	Prod	Oil	Gas	Water
Vessel	dd	mm	m³	10³ m³	m³	hours	m³/hr	10³ m³/hr	m³/hr	hours	m³	10³ m³	m³
Prior mo.	25	6	9.05	1.35	3.53	24.00	0.3771	0.0563	0.1471	96	36.2	5.4	14.1
	5	7	8.85	1.28	3.26	24.00	0.3688	0.0533	0.1358	168	62.0	9.0	22.8
	12	7	9.40	1.51	2.98	24.00	0.3917	0.0629	0.1242	216	84.6	13.6	26.8
	21	7	9.15	1.67	3.65	24.00	0.3813	0.0696	0.1521	264	100.7	18.4	40.2
										Totals			
										744	283.5	46.4	103.9

UID	AB WI 100080100101W400						Hourly test rate				Estimated production		
	Test date		Test oil	Test gas	Test water	Test duration[c]	Oil	Gas	Water	Prod	Oil	Gas	Water
Vessel	dd	mm	m³	10³ m³	m³	hours	m³/hr	10³ m³/hr	m³/hr	hours	m³	10³ m³	m³
Prior mo.	28	6	5.05	0.95	4.15	24.00	0.2104	0.0396	0.1729	48	10.1	1.9	8.3
	3[a]	7	5.85	1.25	4.50	48.00	0.2406	0.0490	0.1792	336	80.8	16.5	60.2
	4[a]	7	5.70	1.10	4.10								
	17	7	6.01	1.15	5.00	25.50	0.2357	0.0451	0.1961	168	39.6	7.6	32.9
	24	7	5.40	0.99	4.10	22.75	0.2374	0.0435	0.1802	192	45.6	8.4	34.6
										Totals			
										744	176.1	34.4	136.0

UID	AB WI 100160100101W400						Hourly test rate				Estimated production		
	Test date		Test oil	Test gas	Test water	Test duration[c]	Oil	Gas	Water	Prod	Oil	Gas	Water
Vessel	dd	mm	m³	10³ m³	m³	hours	m³/hr	10³ m³/hr	m³/hr	hours	m³	10³ m³	m³
Prior mo.	1[b]	7	1.80	1.10	2.20	24.00	0.0750	0.0458	0.0917	24	1.8	1.1	2.2
	2[b]	7	4.00	2.00	5.00	24.00	0.1667	0.0833	0.2083	120	20.0	10.0	25.0
	7	7	3.95	1.95	4.95	23.00	0.1717	0.0848	0.2152	288	49.4	24.4	62.0
	19	7	4.25	2.05	5.05	26.00	0.1635	0.0788	0.1942	216	35.3	17.0	41.9
	28	7	5.65	2.00	5.50	27.75	0.2036	0.0721	0.1982	96	19.5	6.9	19.0
										Totals			
										744	126.0	59.4	150.1

Note that test gas volumes must include gas-in-solution (GIS) volumes (see Section 4.3.5).

[a] Tests on July 3 and 4 were comparable and consecutive (i.e., there were no operational changes). Therefore, the results are combined and used as one 48-hour test.

[b] Tests on July 1 and 2 were not comparable due to operational changes (e.g., choke/pump speed). Therefore, they are used as separate 24-hour tests.

[c] Test duration must be reported to the nearest quarter hour as the minimum resolution (record hours to two decimal places, e.g., 2 hr and 45 min are entered as 2.75 hr).

6.5.2 Calculate Proration Factors and Monthly Production

1) Calculate the total estimated battery production for oil, gas, and water:

   Total estimated battery production = Sum of all the wells' total estimated production

2) Calculate the total actual battery production and proration factors for oil, gas, and water:

   For oil and water,

   Total actual battery production = Total monthly disposition + Closing inventory – Opening inventory – Total receipts

   For gas,

   Total actual battery production = Total monthly disposition (sales, fuel, flare, vent) – Total receipts

   Proration factor = Total actual battery production ÷ Total estimated battery production

   The proration factors for oil, gas, and water must be rounded to five decimal places.

   Note that if a GOR is used to estimate the total battery gas production volume in accordance with Section 4.3.5:

   Estimated battery gas production = Actual battery oil production x GOR

   Estimated battery gas production = Actual battery gas production

   Gas proration factor = 1.00000

3) Calculate each well's monthly prorated production volumes for oil, gas, and water:

   Monthly prorated oil volume = Well estimated oil production x Oil proration factor

   Monthly prorated gas volume = Well estimated gas production x Gas proration factor

   Monthly prorated water volume = Well estimated water production x Water proration factor

4) Check that total well production equals total actual battery production for oil, gas, and water. If the volumes are not equal due to rounding, minor adjustments to the monthly volumes may be required.

   Sum of prorated well production = Total actual battery production

6.6 Condensate Receipts at an Oil Battery

If condensate that could be flashed into the gas phase is received at an oil battery, the licensee must choose from the applicable condensate reporting options in Table 5.2.

If condensate received from an external source will be reported as a GEV, that volume must be subtracted from the total monthly battery gas disposition volume to determine the monthly battery gas production volume.

If condensate that could be flashed off into the gas phase is received at an oil battery, the licensee must report it as below:

- If condensate received into the battery is ≤ 2.0 m^3/day and/or $\leq 5\%$ of the battery oil production, the condensate can be reported as all liquid condensate or as a GEV.

- If condensate received into the battery is > 2.0 m³/day and/or > 5% of the battery oil production, the licensee must report flashed condensate as a GEV receipt volume and unflashed condensate as a liquid condensate receipt.

Note that this may also be applicable to other light hydrocarbons delivered into an oil battery.

6.7 Combined (Cascade) Testing

When a prorated oil well has such low gas production that it cannot properly operate test equipment, a licensee may test two wells simultaneously—combined (cascade) test—through the same test separator. In such cases, the following procedure must be followed:

1) Establish oil, gas, and water production volumes for a high gas producing well by testing it individually through the test separator.

2) Conduct a test for both the high gas producing well and a low gas producing well together through the same test separator immediately after testing the high gas producing well, allowing time for stabilization. (The testing sequence may be reversed with the testing of the combined wells first.)

3) The operating condition of both wells must not be changed. If it is, a new set of tests is required.

4) Total test oil, gas, and water volumes determined for the combined (cascade) test minus the test oil, gas, and water volumes for the high gas producing well will be the test volumes for the low gas producing well (see example below).

5) Both wells should have similar S&W percentages. If any of the calculated oil, gas, or water volumes for the low gas producing well are negative, the tests are not valid and both tests must be repeated.

The use of combined (cascade) testing does not require special approval from the ERCB.

Example
Well A = High gas producing
Well B = Low gas producing

Test Results

Well	Test date	Oil (m³)	Gas (10³ m³)	Water (m³)
Well A + B	July 4	80.0	20.0	20.0
Well A	July 5	50.0	19.0	12.0
Well B = (Well A + B - Well A)	July 4	30.0	1.0	8.0

6.8 Electronic Flow Measurement for Oil Systems

An EFM system is defined as any flow measurement and related system that collects data and performs flow calculations electronically. If it is part of a Distributed Control System (DCS), Supervisory Control and Data Acquisition system (SCADA), or Programmable Logic Controller system (PLC), only the EFM portion has to meet the requirements in this section.

The following systems are not defined as an EFM:

- any meter with an electronic totalizer or pulse counter that does not perform flow calculations (with or without built-in temperature compensation)

- a remote terminal unit (RTU) that transmits any data other than flow data and does not calculate flow

Hardware and software requirements:

- The memory on board the EFM must allow for at least 32 days of storage of the required flow data before being overwritten or erased.

- The EFM must be equipped with its own on-board battery to protect the memory in the event of a power failure.

- The system must have various levels of system security, with the highest level of access to the program restricted to authorized people.

- The communication system must use a data integrity error-checking routine to ensure that the data transmitted are correct.

- The EFM must be set to alarm on high and low differential pressure, if applicable, over-range of any end devices, low power, and communication failures.

- Any changes made to the data or any manually entered values that affect the flow calculation must be flagged so it is clear that these are estimated, not actual, readings.

- This flagging must carry through to values calculated from the data.

6.8.1 Performance Evaluation

If an EFM is used to calculate clean oil volumes, the licensee must be able to verify that it is performing within the ERCB target limits defined in this section.

A performance evaluation test must be completed within **two weeks** after the EFM is put into service and **immediately** after any change to the computer program or algorithms that affects the flow calculation; documentation must be kept and provided to the ERCB upon request. For existing EFM systems, the licensee should conduct its own performance evaluations periodically to ensure that they are performing adequately. A performance evaluation must be conducted and submitted for ERCB audit on request. The ERCB considers either one of the following methods acceptable for performance evaluation:

1) Conduct a performance evaluation test on the system by inputting known values of flow parameters into the EFM to verify the volume calculation and other parameters. The test cases included in this section are for oil/emulsion meters, each with different flow conditions.

   Test cases 1 to 5 are for oil density correction from flowing temperature to 15°C. When a hydrometer is used to measure density, a hydrometer correction is required to compensate for the glass expansion. Density correction to 15°C is only required for blending shrinkage calculations, mass-based measurement, and the CTL and CPL calculations.

   Test cases 6 to 10 are for volume correction using CPL and CTL factors to correct to 101.325 kPa and 15°C. Other manufacturers' recommended methodologies may also be used to evaluate the EFM performance, provided that the volumes obtained from a performance evaluation test agree to within ± 0.1% of those recorded on the sample test cases.

2) Evaluate the EFM calculation accuracy with a flow calculation checking program that performs within the target limits for all the factors and parameters listed in the test cases below. A snapshot of the instantaneous flow parameters and factors, flow rates, and

configuration information is to be taken from the EFM and input into the checking program. If the instantaneous EFM flow parameters, factors, and flow rates are not updated simultaneously, multiple snapshots may have to be taken to provide a representative evaluation.

The densities (test cases 1 to 5) or volumes (test cases 6 to 10) obtained from a performance evaluation test must agree to within ± 0.1% of those recorded on the sample test cases. If the ± 0.1% limit is exceeded, the EFM must be subjected to a detailed review of the calculation algorithm to resolve the deviation problem.

Test Cases for Verification of Oil Flow Calculation Programs

Density and volume corrections in the table below are based on API MPMS, Chapter 11.1 (May 2004).

The hydrometer correction is based on API MPMS, Chapter 9.3 (November 2002).

Density correction to 15°C

	Inputs		Outputs	
	Oil density @ flowing temperature (kg/m³)	Observed temperature (°C)	Oil density corrected to 15°C (kg/m³) (with hydrometer correction)	Oil density corrected to 15°C (kg/m³)(without hydrometer correction)
Test Case 1	875.5	120	942.9	945
Test Case 2	693	11.4	689.9	689.8
Test Case 3	644	84.45	704.7	705.7
Test Case 4	625.5	53.05	660.8	661.4
Test Case 5	779	25	786.7	786.8

Volume correction using pressure and temperature correction factors (CPL and CTL)

	Inputs				Outputs			
	Metered volume (m³)	Oil density @ 15°C (kg/m³)	Flowing temperature (°C)	Flowing pressure (kPa[gauge])	CTL to 15 °C	CPL to 101.325 kPa	CTL corrected volume (m³)	CTL & CPL corrected volume (m³)
Test Case 6	60	903.5	40.5	700	0.98071	1.0005	58.8	58.9
Test Case 7	15	779	3.9	400	1.0112	1.00034	15.2	15.2
Test Case 8	100	1008	89	3700	0.95472	1.00255	95.5	95.7
Test Case 9	250	875.5	5	200	1.00799	1.00013	252	252
Test Case 10	150	640	75	1000	0.90802	1.00365	136.2	136.7

EFM Reports

When any parameter that affects the flow calculation is changed, such as meter factor, fluid densities, or transmitter range, a signoff procedure or an event log must be set up to ensure that the change is made in the EFM system. All data and reports must be retained for a minimum of 12 months.

The required information on each report must be stored using electronic/magnetic (not necessarily on the EFM) or printed media. They may exist individually on different formats or reports, and must be generated upon request by the ERCB, as follows:

- **Daily** for daily report required data
- **Monthly** for monthly report required data

- Event and alarm logs at regular intervals before information is overwritten
- Meter reports on request for audit

The Daily Report

The daily report must include

- meter identification
- daily accumulated flow with indicating flags for estimated flow made by the system or manual inputs and alarms that have occurred for overranging of end devices
- hours on production or hours of flow (specify)
- flow data audit trail—include at least one of the following:
  - instantaneous values for flow rate, operating pressure (if applicable), and temperature taken at the same time each day,
  - average daily values for volumes, operating pressure (if applicable), and temperature, or
  - hourly accumulated flow rate and average hourly values for operating pressure (if applicable) and temperature.

Existing EFM systems that predate *Guide 34* approvals, do not have any of the above audit trail capabilities, and cannot develop the capability due to system limitations should be evaluated for upgrading when new production is tied into the facilities. The ERCB may request upgrades if audit/inspection results indicate that they are warranted.

The Monthly Report

The monthly report is for the entire system, providing data for each measurement point. It is to contain the following at each measurement point as applicable:

- monthly cumulative flow
- flags indicating any change made to flow volumes
- total hours on production or hours of flow (specify)

The Meter Report

The meter report details the configuration of each meter and flow calculation information. These values are used as part of the "audit trail" to confirm that the flow calculation is functioning correctly. The meter report must include the following as applicable and be produced upon request:

1) Instantaneous flow data:
   - instantaneous flow rate
   - instantaneous operating pressure
   - instantaneous flowing temperature
   - CTL
   - CPL

2) Current configuration information:
   - meter identification

- date and time
- atmospheric pressure
- pressure base
- temperature base
- calibrated operating pressure range
- calibrated temperature range
- meter factor and/or k factor
- shrinkage factor

The Event Log

The event log is used to note and record exceptions and changes to the flow parameter, configuration, programming, and database affecting flow calculations, such as

- transmitter range changes
- algorithm changes
- meter factor or k-factor changes
- other manual inputs

The Alarm Log

The alarm log includes any alarms that may have an effect on the measurement accuracy of the system. The time of each alarm condition and the time each alarm is cleared must be recorded. Alarms that must be reported include

- master terminal unit failures
- remote terminal unit failures
- communication failures
- low-power warning
- high/low volumetric flow rate
- overranging of end devices

7 Gas Proration Batteries

This section presents the requirements and exceptions relating to measurement, accounting, and reporting for gas proration batteries.

Gas well operators have the option of not measuring the gas and/or separated liquids at each well site. If the gas and liquids are not separated and measured, they can be prorated. Operators that decide to install prorated systems in accordance with the provisions of this section are accepting higher uncertainty at the wellhead, offset by lower capital and operating costs.

Prorated wells are tested periodically to determine the typical flow rate. The gas and liquids from a number of wells are measured at a group meter, and the volume at the group meter is prorated back to the individual wells based on the most recent test and the hours on stream.

The measurement uncertainty assigned to individual wells within gas proration batteries is greater than for wells where the gas is separated and measured. For this reason, operators should understand the impact of this type of measurement when dealing with partners and third parties.

Prorated wells can be tied in to the same system as measured wells but under separate battery codes. In these cases, the measured wells are kept whole, and the difference between the battery disposition and the measured well volume is prorated to all the proration wells. This is referred to as measurement by difference (see Section 5.5), since the measured volume is subtracted from the group measurement before proration. Measurement by difference increases the uncertainty of the prorated well volume estimate.

7.1 General Requirements

The three types of gas proration batteries allowed in Alberta are

- gas multiwell proration SE Alberta batteries (PRA subtype 363),
- gas multiwell proration outside SE Alberta batteries (PRA subtype 364), and
- gas multiwell effluent measurement batteries (PRA subtype 362).

All wells in a gas proration battery must be classified as gas wells and must be connected by flow line to a common group separation and measurement point.

All gas proration batteries require periodic well tests to be conducted to determine production rates, production ratios, and/or ECF that will be used in the determination of monthly estimated well production volumes. Monthly estimated well production volumes are multiplied by proration factors to determine the actual well production volumes for reporting purposes. All wells must be tested annually unless otherwise stated in this section.

Test taps must be installed at all proration gas wells. The required test tap locations are specified below for each of the above batteries.

See Section 8 of this directive for sampling and analysis of gas, condensate, and water.

7.1.1 Group Measurement

All gas proration batteries require continuous single-phase measurement of total monthly battery gas, condensate, and water production volumes at the group measurement point. If

liquid condensate is trucked out of the group separation and measurement point to a gas plant for further processing, the condensate must be reported as a GEV from the battery, through the gas gathering system, to the gas plant.

Gas wells in any one of the three types of proration batteries must not be commingled with "measured" gas sources or gas from another proration battery without group measurement, or with gas wells in a different type of gas proration battery, upstream of their respective group measurement points. Variances from this requirement may be allowed if the "Exception" criteria in Section 5.5 are met or if site-specific approval has been obtained from the ERCB prior to implementation.

7.1.2 Stabilized Flow and Representative Flow

The words stabilized flow and representative flow are used extensively in this section as defined below.

Stabilized flow indicates a point at which flowing parameters of gas, condensate, or water are producing under normal operating conditions and represent production levels equal to the well's normal average flow rate. Stabilized flow can only be achieved when all testing equipment associated in determining an actual volume has reached equilibrium (i.e., liquid levels in test separator, pressure and temperature stabilization to normal operating conditions).

Representative flow can be used when stabilized flow is not achievable, such as for wells with artificial lift systems and wells with slugging characteristics. The test volumes of gas, condensate, or water must be representative of the well's production capability under normal operating conditions.

Wells that use artificial lift systems or characteristically display slug flow must be tested for a minimum duration that completes multiple flow cycles to accurately determine a representative volume of gas, condensate, or water. These representative production volumes are then extrapolated to accurately reflect the wells' production over an extended period of time.

7.1.3 Required Decimal Places for Volumetric Calculations in Gas Proration Batteries

Type of calculations	Number of decimals to be calculated to	Number of decimals to be rounded to
Productions and estimated productions	2	1
Well test gas, GEV of test condensate, test condensate, or test water	3	2
WGR, condensate-gas ratio (CGR), and oil-gas ratio (OGR)	5	4
Proration factors, ECF	6	5

7.2 Gas Multiwell Proration SE Alberta Batteries

Gas wells in this type of battery do not require dedicated continuous measurement for each well or special approvals from the ERCB.

Production rates determined during a well test must be used in the estimation/proration calculations within 30 days of the test until the next test is conducted.

Total battery gas production must be measured and prorated back to the individual wells, based on each well's estimated monthly gas production. Estimated well gas production is based on hourly production rates, determined by periodic well tests and monthly producing hours.

Gas wells that produce from shallow gas zones in SE Alberta may be included in these types of batteries. The zones may include, but are not limited to, the Medicine Hat, Milk River, Second White Specks, Belly River, and Colorado zones. The production from two or more of these zones without segregation in the wellbore requires either prior approval from the ERCB for commingled production, which has been granted in a portion of SE Alberta in Order No. MU 3847, or adherence to the self-declared commingled production requirements described in *Directive 065: Resources Applications for Conventional Oil and Gas Reservoirs.*

7.2.1 Group Measurement

Group measured production is generally determined through individually measured product streams. A minimum of two-phase group measurement is required because the battery water production must be reported at the battery level. This group measurement point is located generally at the battery site where a compressor is present (see Figure 7.1).
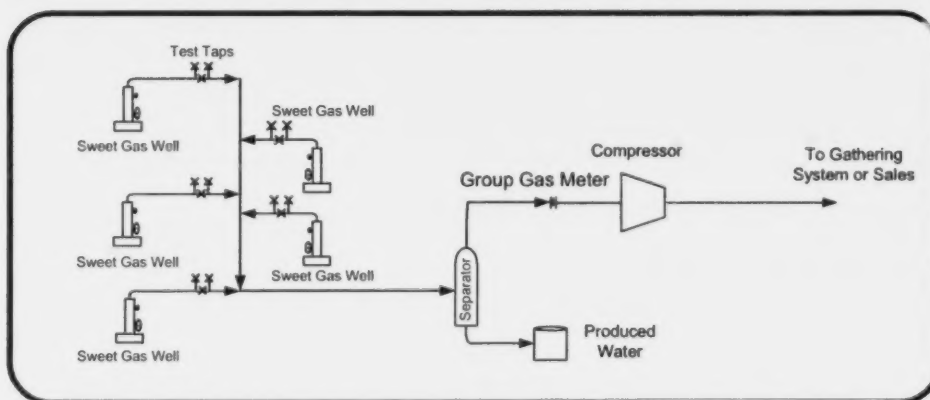
Figure 7.1. Typical SE Alberta shallow gas battery

7.2.2 Size of a SE Alberta Multiwell Gas Proration Battery

There is no limit on the number of flow-lined wells that may be in a SE Alberta shallow gas proration battery; however, licensees are encouraged to consider the logistics of the battery's operation in determining the size of these batteries, with the key factors being

- the ability to conduct representative well tests at the minimum frequency specified below, and

- the configuration and operating pressures of the battery and flow lines such that all wells can readily flow.

This approach will generally result in the main pipeline system laterals being used to establish a group measurement point.

7.2.3 Testing Requirements

Gas production rate tests must be conducted for each well in the battery in accordance with the following requirements:

- The test must be of sufficient duration to clearly establish a stabilized flow rate.

- The test must be representative of the well's capability under normal operating conditions.

- Testing programs and procedures must ensure that all wells are treated equitably within their respective batteries. These types of wells are typically tested by directing flow from the well through a test meter; however, a test separator system could also be used.

- New wells must be tested within the first 30 days of production, then again within 12 months, and thereafter according to Table 7.1.

If these requirements cannot be satisfied, the operator must either reconfigure the system (e.g., redirect some wells to another battery/group measurement point) or test each of the individual wells within the battery once per month.

Table 7.1. Testing frequency for SE Alberta shallow gas wells

te	ate	tests	
	d		
d	d		
d			

*See Section 2.3 for frequency definition.

7.2.4 Production Accounting and Reporting Procedures

Water Reporting Requirements

The reporting of water production for the qualified wells in SE Alberta shallow gas batteries is not required. However, all water receipts and disposition must be reported at the battery level. The water must be properly stored in accordance with ERCB *Directive 055* requirements. An "ABMC" receipt code can be used to balance the disposition at the battery level on the PRA. If the water is trucked to non-PRA reporting facilities without a reporting code or evaporated on site, it must be reported using an "ABMC" disposition code.

Production Volume Calculations

Monthly gas production volumes are to be calculated as follows:

1) Calculate well gas test rate:

Well gas test rate $(10^3 \text{ m}^3/\text{hour})$ = Well test gas volume (10^3 m^3) / Well test duration (hours)

2) Calculate estimated monthly well gas volume:

Estimated monthly well gas volume = Well gas test rate x Monthly total hours of well production

3) Calculate total estimated gas production for the battery:

Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

4) Calculate proration factor for gas:

Gas proration factor = Total battery measured monthly gas volume / Total battery estimated monthly gas volume

5) Calculate actual monthly (prorated) well gas production:

Actual monthly well gas production = Gas proration factor x Estimated monthly well gas volume

7.3 Gas Multiwell Proration Outside SE Alberta Batteries

Gas wells in this type of battery do not require dedicated continuous measurement for each well. Production rates, WGRs, and/or CGRs determined during a well test must be used as soon as reasonably possible in the estimation/proration calculations until the next test is conducted.

Total battery gas production must be measured and prorated back to the individual wells based on each well's estimated monthly gas production. Estimated well gas production is based on hourly production rates, determined by periodic well tests and monthly producing hours.

Total battery condensate production must be measured if present. If it is delivered for sale from the battery, it must be prorated back to the individual wells based on each well's CGR from the production tests. The sales condensate must be reported as a liquid disposition on the PRA. Then the estimated gas production volume at each well will not include the GEV of the condensate. If the condensate is recombined with the gas for further processing to a gas plant, the condensate must be reported as a GEV and added to the measured gas production volume and reported on the PRA.

Total battery water production must be measured and prorated back to the individual wells based on each well's estimated monthly water production. Estimated well water production is based on a WGR, determined by periodic well tests multiplied by the estimated monthly well gas production (see Figure 7.2).
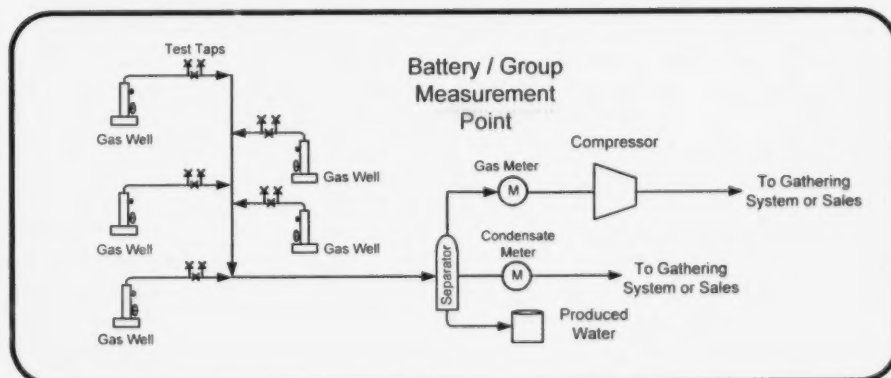
Figure 7.2. Typical outside SE Alberta multiwell gas proration battery

If total water production at each well in the battery is less than or equal to 0.50 m³/d based on the monthly average flow rates recorded during the six months prior to conversion, water production may be prorated to all wells in the battery based on the estimated gas production

at each well. If a group of new wells not previously on production is to be configured as a proration battery, the qualifying flow rates must be based on production tests conducted under the normal operating conditions of the proration battery.

There is no geographical or zonal limitation for this type of proration battery. The "Exception" criteria in Section 5.4 must be met or ERCB site-specific approval must be obtained prior to the proration battery implementation either at the initial design and installation stage or at a later stage of production when the production rate decreases to a point that continuous measurement is not economical.

Gas wells classified as producing oil rather than condensate must not be tied into a gas multiwell proration outside SE Alberta battery unless the well oil and gas production volumes are separated and measured prior to commingling with the other wells in the battery and either the "Exception" criteria in Section 5.5 are met or site-specific approval has been obtained from the ERCB prior to implementation. However, if a gas well classified as producing condensate in a gas multiwell proration outside SE Alberta battery is reclassified by the ERCB as producing oil, the well may remain in the battery provided that the well is equipped with a separator and there is continuous measurement of the gas, oil, and water or, alternatively, the measurement, accounting, and reporting procedures specified in Section 7.3.2.1 are followed.

7.3.1   Well Testing Requirements

Well testing is typically performed by directing well production through a three-phase portable test separator configured with dedicated meters for gas, condensate, and water. Test equipment using two-phase separation is acceptable if hydrocarbon liquids are too small to be measured during typical well test durations. Other options that provide equivalent liquid volume determination accuracy may also be considered. For example, if a three-phase separator is not available, alternative equipment, such as a two-phase separator with a total liquid meter and continuous water cut analyzer, may be acceptable (see Figures 7.3 and 7.4).
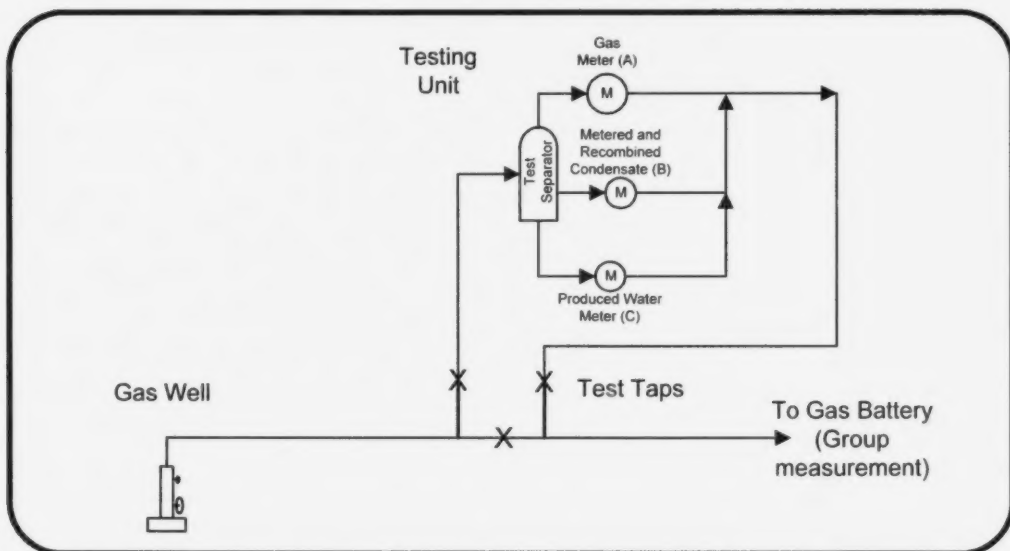
Figure 7.3. Typical testing unit for gas multiwell proration outside SE Alberta battery
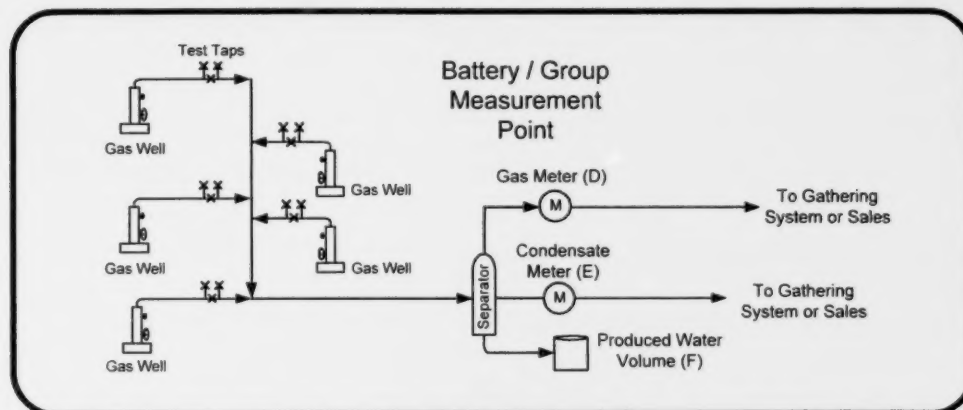
Figure 7.4. Typical gas multiwell proration outside SE Alberta battery

Test frequency may be extended with ERCB approval.

Unless alternative test procedures have been specified in an ERCB approval, the test must be conducted with measurement of all phases as follows:

- The test must begin only after a liquid level stabilization period.

- The test duration must be a minimum of 12 hours.

- After the commencement of production at the proration battery, all wells must be tested within the first month, then again within six months, and thereafter annually. New wells added to the battery at some future date must be tested within the first month of production, then again within six months, and thereafter annually.

- Consistent testing procedures must be used for consecutive tests to identify if a change in a well's flow characteristics has occurred.

- These wells are typically tested by directing flow from the well through a test separator. If the initial testing with a separator shows a liquid-gas ratio (LGR) of less than 0.01 m^3 liquid/10^3 m^3 gas, other testing methodology, such as a smaller separator or a single test meter without separation, could be used for the next test. If the total liquid volumes at group measurement point exceed a ratio of 0.05 m^3 liquid/10^3 m^3 gas in any month, a test separator must be used to test all the wells within the battery for the next round of testing to determine where the liquid originated.

- The gas, condensate, and water volumes must be measured.

- The condensate must be sampled during every test and subjected to a compositional analysis, which is to be used to determine the gas equivalent factor (GEF). The sample may be taken from the condensate leg of a three-phase separator or the liquid leg of a two-phase separator. (The water must be removed from the condensate before conducting the analysis.)

- The GEF must be used to convert the liquid condensate volume determined during the test to a GEV, which will be added to the measured test gas volume to determine the total test gas volume if the condensate is not delivered for sale at the group measurement point (see Section 7.3.2).

- The WGR, CGR, and OGR (if applicable) must be determined by dividing the test water, condensate, and oil volume respectively by the total test gas volume.

- For orifice meters, the test gas meter must use 24-hour charts for a test period of 72 hours or less, unless electronic flow measurement is used; for testing periods longer than 72 hours, 7-day charts may be used, provided that good, readable pen traces are maintained (see Section 4.3.4).

7.3.2 Production Volume Calculations

Monthly production volumes are to be calculated as follows:

Units: All gas volumes and GEV are to be in (10^3 m^3) and liquid volumes in (m^3).

1) Calculate well gas test rate (see Figure 7.3):

    Well gas test rate $(10^3 \text{ m}^3/\text{hour})$ = (Well test gas volume [A] + GEV of well test condensate [B]) / Well test hours

    Note: Do not include GEV of [B] if condensate is delivered for sale at the group measurement point.

2) Calculate estimated monthly well gas volume:

    Estimated monthly well gas volume = Well gas test rate x Monthly total hours of well production

3) Calculate total estimated gas production for the battery:

    Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

4) Calculate the well WGR (see Figure 7.3):

    WGR = Well test water volume (C) / (Well test gas volume [A] + GEV of well test condensate [B])

    Note: Do not include GEV of [B] if condensate is delivered for sale at the group measurement point.

5) Calculate estimated water production for each well:

    Estimated monthly well water volume = Estimated monthly well gas volume x WGR

6) Calculate total estimated water production for the battery:

    Total battery estimated monthly water volume = Sum of all estimated monthly well water volumes

If the condensate is delivered for sale at the group measurement point, calculate the next two items; otherwise go directly to item 9.

7) Calculate the well CGR (see Figure 7.3):

    CGR = Well test condensate volume (B) / Well test gas volume (A)

8) Calculate estimated condensate production for each well:

    Estimated monthly well condensate volume = Estimated monthly well gas volume x CGR

9) Calculate total estimated condensate production for the battery:

    Total battery estimated monthly condensate volume = Sum of all estimated monthly well condensate volumes

10) Calculate proration factors for gas, condensate (if delivered for sale), and water (see Figure 7.4):

Gas Proration Factor (GPF) = (Total battery measured monthly gas volume [D] + GEV of total battery condensate [E]) / Total battery estimated monthly gas volume

Note: Do not include GEV of [E] if condensate is delivered for sale at the group measurement point.

Water Proration Factor (WPF) = Total battery actual monthly water volume (F) / Total battery estimated monthly water volume

Condensate Proration Factor (CPF) = Total battery measured monthly condensate volume [E] / Total battery estimated monthly condensate volume

11) Calculate actual monthly (prorated) well production:

Actual monthly well gas production = Estimated monthly well gas volume x GPF

Actual monthly well water production = Estimated monthly well water volume x WPF

Actual monthly well condensate production = Estimated monthly well condensate volume x CPF

7.3.3 Exception

If a gas well classified as producing condensate in a gas multiwell proration outside SE Alberta battery is reclassified by the ERCB as a gas well producing oil, the well may remain in the battery provided that the well is equipped with a separator and there is continuous measurement of the gas and liquid components or, alternatively, the measurement, accounting, and reporting procedures specified below are followed (see Figure 7.5).

Annual Gas Rate–WGR tests must be conducted on the well. An oil-gas ratio (OGR) must also be determined during this test. The WGR, estimated water production, water proration factor, and actual water production are determined in the same manner as indicated above (see Section 7.3.2).

Units: All gas volumes and GEV are to be in (10^3 m^3) and liquid volumes in (m^3).

1) Calculate well gas test rate (see Figure 7.6):

Well gas test rate = Well test gas volume (A) / Well test hours

2) Calculate estimated monthly well gas volume:

Estimated monthly well gas volume = Well gas test rate x Monthly total hours of well production

3) Calculate the OGR (see Figure 7.6):

OGR = Well test oil volume (B) / Well test gas volume (A)

4) Calculate actual well oil production:

Actual monthly well oil production = Estimated monthly well gas volume x OGR

5) Calculate actual total oil production:

Actual monthly total battery oil production = Sum of all actual monthly well oil volume

6) At the group measurement point, subtract the oil production volume (item 5) from the total liquid hydrocarbon volume to determine the total battery condensate production (the GEV of the total battery condensate volume, if not delivered for sale, must be added to

the measured group gas volume to determine the total battery gas volume) (see Figure 7.5):

Total battery condensate volume = Battery total liquid hydrocarbon volume (E) –
Actual monthly total battery oil production

7) Calculate total estimated gas production for the battery:

Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

8) Calculate proration factor for gas (see Figure 7.5):

Gas Proration Factor (GPF) = (Total battery measured monthly gas volume [D] +
GEV of total battery condensate volume [item 6]) /
Total battery estimated monthly gas volume

9) Calculate actual monthly (prorated) well gas production:

Actual monthly well gas production = Estimated monthly well gas volume x GPF
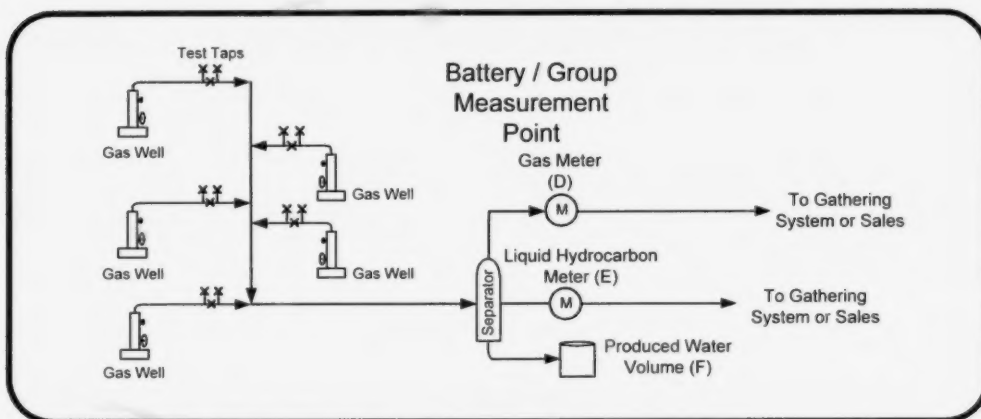
Figure 7.5. Typical gas multiwell proration outside SE Alberta battery
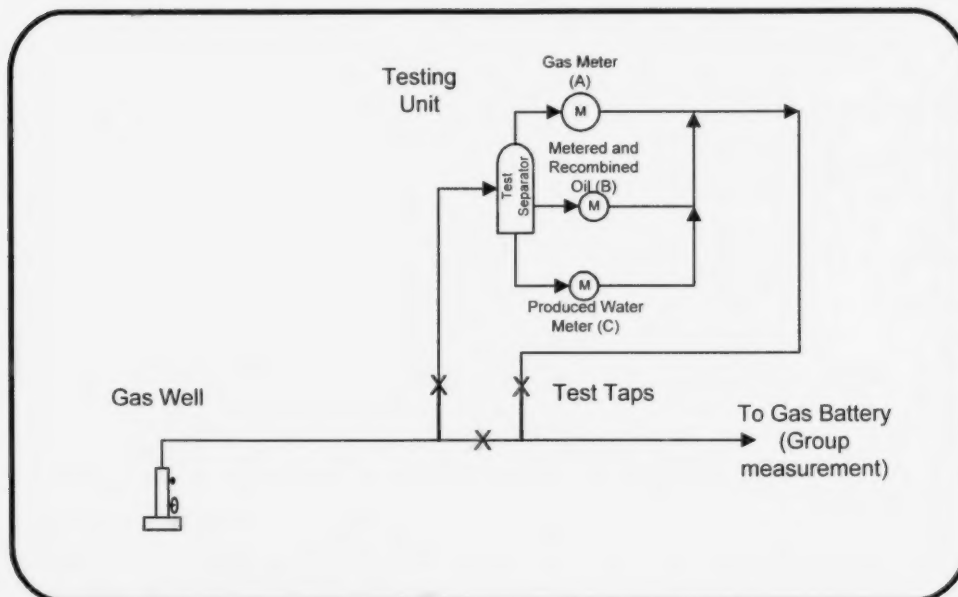
Figure 7.6. Typical testing setup

Report the calculated monthly oil production volume as oil produced from the well. Prorate monthly condensate (if delivered for sale) and water production as in the normal proration battery in Section 7.3.2.

7.4 Gas Multiwell Effluent Proration Batteries

Gas wells in this type of battery have dedicated "effluent" or "wet gas" measurement, whereby total multiphase well fluid passes through a single meter (see Figure 7.7). This type of measurement must be subjected to testing regardless of the type of effluent meter used.
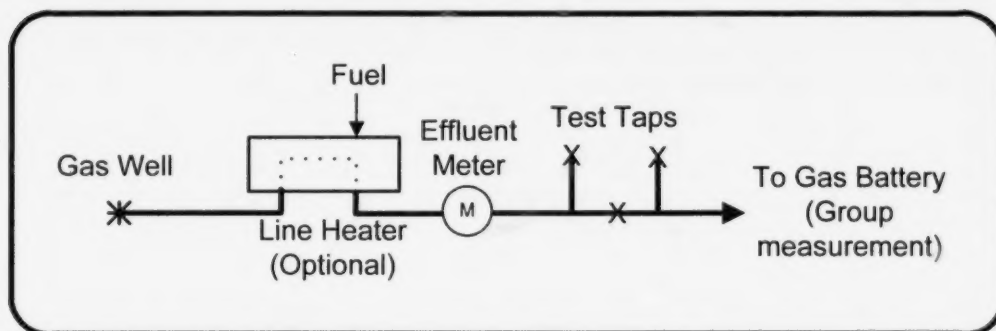
Figure 7.7. Typical gas well effluent metering configuration

Production rates, WGR, CGR, and ECF determined during a well test must be used in the estimation/proration calculations within 60 days of the test until the next test is conducted.

Total battery gas production must be measured and prorated back to the individual wells, based on each well's estimated monthly gas production. Estimated well gas production is based on the total volume measured by the effluent meter multiplied by an ECF (see Figure 7.8).

For most wells, the required minimum well testing frequency is annual unless the criteria in Section 5.7 are met. Total battery water production must be measured and prorated back to the individual wells, based on each well's estimated monthly water production. Estimated well water production is based on a WGR, determined by periodic well tests multiplied by the estimated monthly well gas production.

Generally, a total liquid rate of less than or equal to 0.28 m^3 liquid per 10^3 m^3 gas is recommended for this type of metering application. The uncertainty of measurement will increase with higher liquid rates, especially under liquid slugging conditions. However, there are no liquid rate limits to effluent measurement.

Gas wells that are classified as producing oil, rather than condensate, must not be tied into an effluent proration battery, unless the well oil and gas production volumes are separated and measured prior to commingling with the effluent wells and either the "Exception" criteria in Section 5.5 are met or site-specific approval has been obtained from the ERCB prior to implementation. However, if a gas well classified as producing condensate in a multiwell effluent proration battery is reclassified by the ERCB as producing oil, the well may remain in the battery provided that the well is equipped with a separator and there is continuous measurement of the gas and liquid components or, alternatively, the measurement, accounting, and reporting procedures specified in Section 7.4.3 are followed.
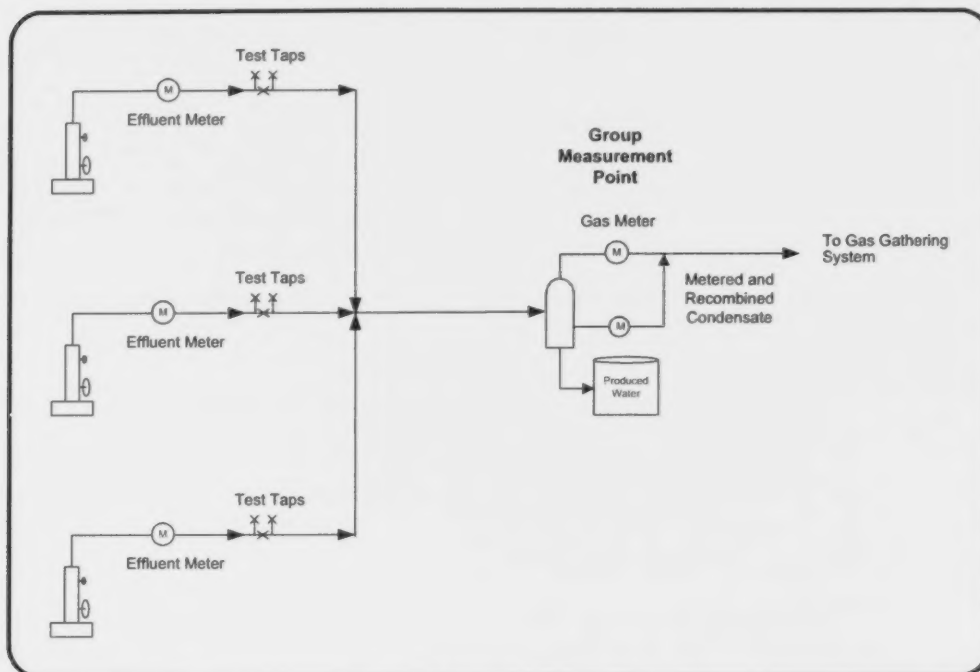
Figure 7.8. Typical multiwell effluent proration battery configuration

Figure 7.7 illustrates a typical gas well effluent measurement configuration. Production from the gas well passes through a line heater (optional), where it is heated. This is typically done to vapourize some of the hydrocarbon liquids and heat up the water and the gas in the stream before metering to prevent hydrate formation. For well testing purposes, test taps must be located downstream of this meter within the same pipe run. The line heater and/or the fuel gas tap, if present, must be upstream of the meter or downstream of the test taps to ensure that the test measurement is subjected to the same condition as the effluent meter. After measurement, production from the well is commingled with other flow-lined effluent gas wells in the battery and sent to a group location, where single-phase (group) measurements of hydrocarbon liquids, gas, and water must be conducted downstream of separation.

7.4.1 Well Testing

Well testing is performed by directing well production downstream of the effluent meter and within the same pipe run through a three-phase portable test separator configured with dedicated meters for gas, condensate, and water (see Figure 7.9). Test equipment using two-phase separation is acceptable if hydrocarbon liquids are too small to be measured during typical well test durations. Other options that provide equivalent liquid volume determination accuracy may also be considered. For example, if a three-phase separator is not available, alternative equipment, such as a two-phase separator with a total liquid meter and continuous water cut analyzer, may be acceptable. The test must be conducted as follows:

- The test must begin only after a liquid level stabilization period within the test separator.

- The test duration must be a minimum of 12 hours.

- The required minimum well testing frequency is annual unless the criteria specified in Section 5.7 are met. All new wells must be tested within the first 30 days of initial production.

- Consistent testing procedures must be used for consecutive tests to identify if a change in a well's flow characteristics has occurred.

- The gas, condensate, and water volumes must be measured.

- The condensate must be sampled during every test and subjected to a compositional analysis, which is to be used to determine the GEF. The sample may be taken from the condensate leg of a three-phase separator or the liquid leg of a two-phase separator (the water must be removed from the condensate before conducting the analysis).

- The GEF must be used to convert the liquid condensate volume determined during the test to a GEV, which will be added to the measured test gas volume to determine the total test gas volume if the condensate is not delivered for sale at the group measurement point (see Section 7.4.2).

- The WGR must be determined by dividing the test water volume by the sum of the measured test gas volume and the gas equivalent of the measured test condensate volume if the condensate is not delivered for sale at the group measurement point (see Section 7.4.2).
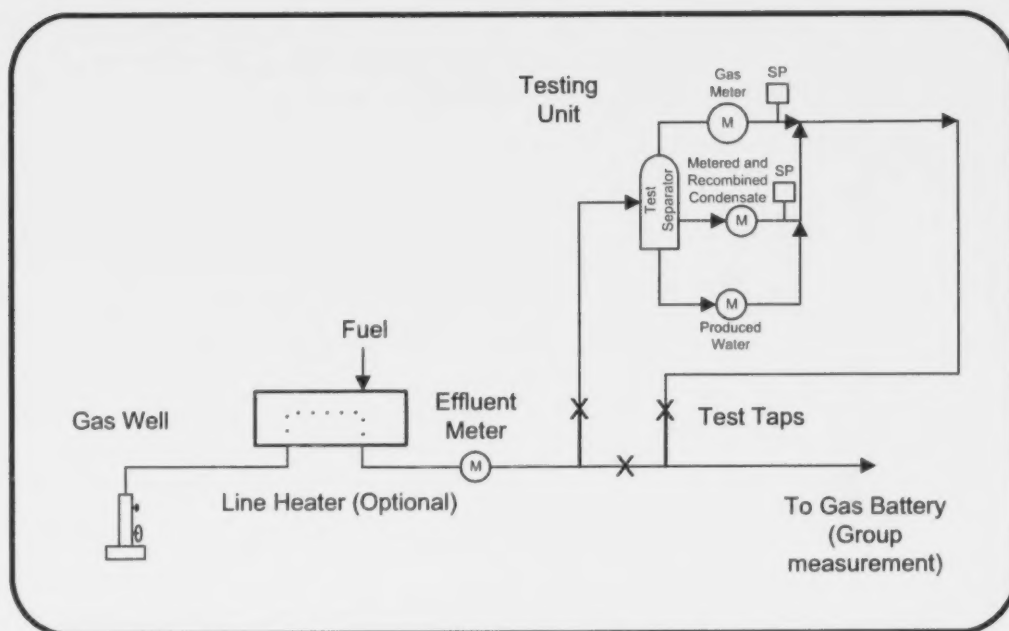
Figure 7.9. Typical effluent well measurement configuration with test unit

- The ECF must be determined by dividing the sum of the measured test gas volume and the GEV of the measured test condensate volume by the effluent metered gas volume.

- For orifice meters, the effluent meter and the test gas meter must use 24-hour charts for a test period of 24 hours or less, unless electronic flow measurement is used; for testing periods longer than 24 hours, 7-day charts may be used, provided that good, readable pen traces are maintained (see Section 4.3.4).

7.4.2 Production Volume Calculations

Monthly production volumes are to be calculated as follows (see Figure 7.10):

Units: All gas volumes and GEV are to be in (10^3 m^3) and liquid volumes in (m^3).

If condensate is delivered for sale at the group measurement point, go directly to item 8; otherwise follow items 1 to 7:

1) Calculate the ECF:

   ECF = (Well test gas volume [B] + GEV of well test condensate [C]) /
         Effluent gas volume measured during test (A)

2) Calculate estimated gas production for each well:

   Estimated monthly well gas volume = Monthly well effluent volume x ECF

3) Calculate the WGR:

   WGR = Well test water volume (D) / (Well test gas volume [B] + GEV of well test condensate [C])

4) Calculate estimated water production for each well:

   Estimated monthly well water volume = Estimated monthly well gas volume x WGR
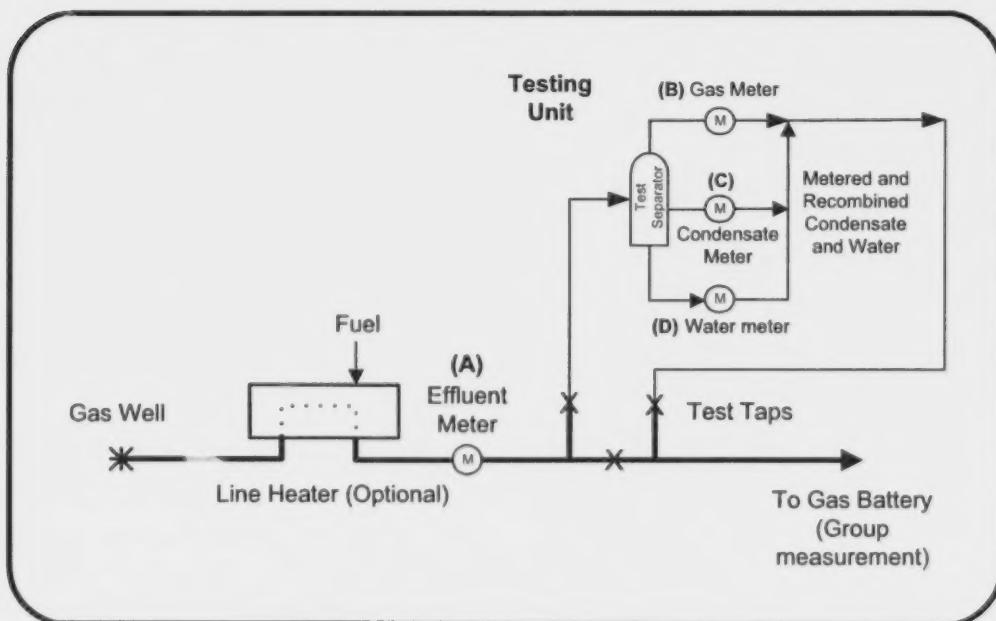
Figure 7.10. Effluent well meter testing configuration with condensate production

5) Calculate total battery estimated volumes (gas and water):

   Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

   Total battery estimated monthly water volume = Sum of all estimated monthly well water volumes

6) Calculate proration factors for gas and water:

GPF = (Total battery measured monthly gas volume + GEV of total battery condensate) /
Total battery estimated monthly gas volume

WPF = Total battery actual monthly water volume / Total battery estimated monthly water volume

7) Calculate actual monthly (prorated) well production:

Actual monthly well gas production = Estimated monthly well gas volume x GPF

Actual monthly well water production = Estimated monthly well water volume x WPF

For the battery with condensate delivered for sale at the group measurement point:

8) Calculate the ECF:

ECF = Well test gas volume [B] / Effluent gas volume measured during test (A)

9) Calculate the well CGR:

CGR = Well test condensate volume (C) / Well test gas volume (B)

10) Calculate the WGR:

WGR = Well test water volume (D) / Well test gas volume [B]

11) Calculate estimated gas, condensate, and water production for each well:

Estimated monthly well gas volume = Monthly well effluent volume x ECF

Estimated monthly well condensate volume = Estimated monthly well gas volume x CGR

Estimated monthly well water volume = Estimated monthly well gas volume x WGR

12) Calculate total estimated gas, condensate, and water production for the battery:

Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

Total battery estimated monthly water volume = Sum of all estimated monthly well water volumes

Total battery estimated monthly condensate volume = Sum of all estimated monthly well condensate volumes

13) Calculate proration factors for gas, condensate, and water:

Gas Proration Factor (GPF) = Total battery measured monthly gas volume / Total battery estimated monthly gas volume

Condensate Proration Factor (CPF) = Total battery measured monthly condensate volume / Total battery estimated monthly condensate volume

Water Proration Factor (WPF) = Total battery actual monthly water volume / Total battery estimated monthly water volume

14) Calculate actual monthly (prorated) well production:

Actual monthly well gas production = Estimated monthly well gas volume x GPF

Actual monthly well condensate production = Estimated monthly well condensate volume x CPF

Actual monthly well water production = Estimated monthly well water volume x WPF

7.4.3   Exception

If an existing gas well classified as producing condensate in a multiwell effluent proration battery is reclassified by the ERCB as a gas well producing oil, the well may remain in the multiwell effluent proration battery provided that the well is equipped with a separator and there is continuous measurement of the gas and liquid components or, alternatively, the effluent meter is left in place and the measurement, accounting, and reporting procedures specified below are followed (see Figure 7.11).

Annual ECF-WGR tests must be conducted on the well. (These types of wells do not qualify for the test frequency exemptions or reductions described above.) An OGR must also be determined during this test. The WGR, estimated water production, water proration factor, and actual water production are determined in the same manner as indicated above.
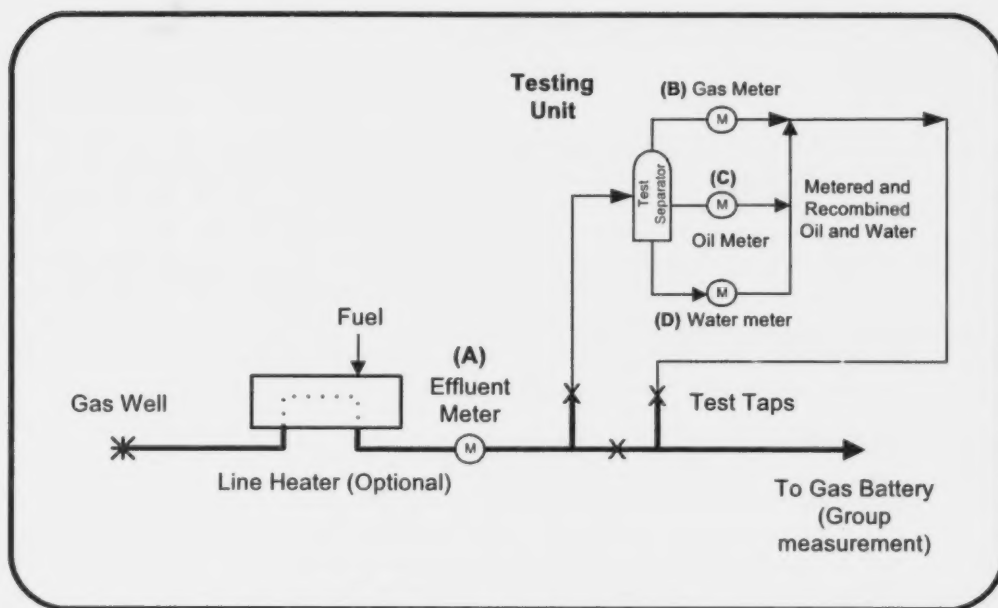
Figure 7.11. Effluent well meter testing configuration with oil production

1) Calculate the ECF:

   ECF = Well test gas volume (B) / Effluent gas volume measured during test (A)

2) Calculate estimated gas production for the well:

   Estimated monthly well gas volume = Monthly well effluent volume x ECF

3) Calculate the OGR:

   OGR = Well test oil volume (C) / Well test gas volume (B)

4) Calculate actual well oil production:

   Actual monthly well oil volume = Estimated monthly well gas production x OGR

5) Calculate the total monthly battery condensate volume:

   Total battery condensate volume = Total battery liquid hydrocarbon volume –
           Total monthly oil volume

6) Report the calculated monthly oil production volume as oil produced from the well. Prorate monthly gas and water production as in Section 7.4.2.

8 Gas and Liquid Sampling and Analysis

Introduction

This section outlines the gas and related liquid sampling and analysis requirements for the various categories of production measurement. These requirements add to the requirements in Sections 11.070 and 11.080 of the *OGCR*, which continue to apply. The requirements vary, depending on a number of factors, such as production rate, potential for the composition to change over time, and the end use of the fluid. Where appropriate, conditions have been identified under which the sampling and analysis requirements may be altered or eliminated altogether. The ERCB will also consider applications for further requirement alterations or eliminations if the licensee can demonstrate that measurement accuracy would either not be reduced or not impact royalty, equity, or reservoir engineering concerns.

8.1 General

Gas and liquid analyses are required for the determination of gas volumes, conversion of liquid volumes to gas equivalent, and product allocation. The sampling and analysis requirements identified in this section pertain only to those areas that affect the calculations and reporting required by the ERCB.

These requirements apply solely to the measurement of hydrocarbon fluids and are not intended to supersede the business requirements that licensees are required to meet regarding product allocations.

If oil is produced from gas wells (as defined by the ERCB), it must be reported as liquid oil production and not as a gas equivalent volume (GEV), so compositional analysis of the oil is not required for that purpose. The oil produced could be combined with the gas and delivered to a gas plant or other facilities for further processing, or the oil could be separated from the gas at the well equipment and directed to tankage, and then on to sales or further treatment.

Gas density and composition are integral components of gas volume calculations and plant product allocation calculations. For differential producing meters, such as orifice meters, venturi meters, and flow nozzles, the accuracy of a computed volume and component allocations are very sensitive to the accuracy of the compositional analysis, which is the basis for compressibility factors and density determination. For linear meters, such as ultrasonic and vortex, the compositional analysis is primarily used to determine the compressibility factors.

If liquid condensate produced from gas wells is either recombined with the gas well production or trucked to the inlet of a gas plant for further processing, the compositional analysis from a condensate sample must be used to determine the GEV of the condensate, which must be added to the well gas volume for reporting purposes. A similar procedure applies to gas gathering systems where liquid condensate is delivered to other facilities for further processing and to gas plant inlets. For this reason, the condensate sampling requirements must mirror the gas sampling requirements.

If liquid condensate is separated at a well, battery, or gas gathering system and delivered from that point for sale or other disposition without further processing, the condensate must be reported as a liquid volume. Therefore, a compositional analysis of the condensate is not required for gas equivalent volume determination purposes but may be required for the purposes of the sale.

Sampling and analysis frequencies and updating requirements for the various production types are summarized in Section 8.4. Further details are provided in the sections that follow. These sampling frequencies are the base requirements for gas and related liquid measurement.

Sampling and analysis of oil/emulsion streams at oil and gas wells and batteries are performed to determine the relative oil and water content of the streams. Oil/emulsion sampling and analysis are discussed in Section 6.4.

8.2   Sampling Requirements

Except where noted in this directive, the gas sampling equipment and methodology must follow the requirements set out in API MPMS 14.1 (June 2001), Gas Processors Association (GPA) 2166-05, or other equivalent industry standards.

Except where noted in this directive, the condensate sampling equipment and methodology must follow the requirements set out in GPA 2174-93, the evacuated cylinder method cited in GPA 2166-05, or in other equivalent industry standards.

Samples and analysis may be obtained by any of the following methods:

- on-site gas chromatograph (GC)
- proportional sampling
- spot or "grab" sampling

Spot or grab samples are acceptable for obtaining gas and liquid analyses once per test or per determination, provided that uncertainty requirements in Section 1 are fulfilled. When the uncertainty requirements cannot be met, licensees must consider more frequent sampling, calculated analyses (see Section 8.3.1), proportional samplers, or chromatographs.

For example: If the analysis from one time period to the next is such that the density and/or compressibility changes cause the volume to change by more than the allowable uncertainty, a more frequent analysis is required or an alternative method of obtaining the sampled must be used.

The gas and liquid analyses must be updated when operating conditions are significantly altered (e.g., addition/removal of compression or line-heating, addition/removal of production sources in a common stream, wellbore recompletion).

If the gas volumes for all meters in the common stream (e.g., sales, fuel, flare, and injection gases) meet the uncertainty guidelines in Section 1, the licensee may use a single gas analysis for all meters on the common stream.

Sampling Procedures

1) Sample points must be located to provide representative samples.

2) Sample probes must not be located within the minimum upstream straight lengths of the meter.

3) Access from grade or platform must be provided for the sample point.

4) If sample transfer tubing is to be used, its length must be minimized.

5) The sample transfer tubing must be oriented to minimize the potential to trap liquids in gas samples and water in condensate samples.

6) A means must be provided to safely purge sample transfer tubing between the sample point and the connection point of the sample cylinder.

7) Sample containers must be clean and meet the pressure, temperature, and materials requirements of the intended service and have the required regulatory approvals as necessary.

8) The procedures used for sampling, transportation, handling, storage, and analysis must ensure that atmospheric contamination does not occur.

The samples must be analyzed using a gas chromatograph or equivalent to determine the components to a minimum of C7+ composition. The gas composition analysis must be determined to a minimum of four decimal points (as a fraction of 1.0000) or two decimal points (as a percentage of 100), and the relative density must be determined to a minimum of three decimal points.

8.2.1 Sample Probes

The sample probe location and installation requirements that follow apply to all ERCB measurement points.

With the exception of sales/delivery (royalty trigger) point measurement, current sample probe locations and installations existing prior to December 5, 2007, do not have to be upgraded to meet the following requirements. A sample probe must be installed according to the requirements below when an installation is relocated or reused for another well or facility.

Requirements for Gas Sampling

1) For sampling applications where the gas is at or near its hydrocarbon dew point, a sample probe must be used (e.g., any separator application where hydrocarbon liquids are present).

2) For gas applications where the gas is not near its hydrocarbon dew point, a sample probe is recommended.

3) The preferred location for gas sample points is the top of horizontal lines.

4) An optional location for gas sample probes is the side of vertical lines with the probe tip sloping 45° downward.

5) Sample probes should be located at least 5 pipe diameters downstream of any piping disturbances, such as bends, elbows, headers, and tees.

6) The location of the sample point must be such that phase changes due to changes in pressure and/or temperature are minimized. Specifically, for gases at or near their hydrocarbon dew point, sample points must not be located downstream of pressure-reducing components, such as control valves, flow conditioners, and regulators, or long lengths of un-insulated piping or within 5 pipe diameter downstream of an orifice plate.

7) Sample points may be located downstream of ultrasonic meters that experience minimum pressure drop through the meter unless a flow conditioner is used and the gas is at or near

its hydrocarbon dew point, in which case the sample point must be upstream of the flow conditioner.

8) Insulation and heat tracing must be used to eliminate any cold "spots" between the sample point and the entry point into the sample container or gas chromatograph where the sample transfer tubing temperature falls below the hydrocarbon dew point, such as at all separator applications.

9) Sample points used to sample blends of two gas streams should have provision for mixing (e.g., upstream static mixer), with due consideration to potential phase changes brought about by a pressure drop associated with the mixing device.

10) Orifice meter impulse lines or transmitter manifolds lines must not be used for taking samples.

11) Level gauge connections must not be used for taking samples.

Requirements for Condensate Sampling

1) With the exception of two-phase separators, a sample probe is recommended.

2) A sample probe must be installed for samples to be used to determine water cut when there is emulsion or a mix of water and hydrocarbon, such as two-phase separators. For such applications, the sampling system design must meet the requirements of API MPMS 8.2 with respect to the use of mixers, sample probe location, and design.

3) The preferred location for condensate sample points is the side of horizontal lines.

4) An optional location for liquid sample points is the side of vertical lines with the probe tip sloping 45° downward.

5) The location of the sample point must be such that phase changes due to changes in pressure and/or temperature are minimized. Specifically, sample points must not be located where vapour breakout is likely (e.g., downstream of pressure-reducing components, such as orifice plates, flow conditioners, turbine, PD or Coriolis mass meters, control valves, and regulators or where the stream temperature has increased).

6) For separator applications, the sample point should be between the separator outlet and the flow/level control valve upstream of the meter, unless a pressure booster pump is used, in which case the sample point is between the pump discharge and the meter.

7) Orifice meter impulse lines or transmitter manifolds lines must not be used for taking samples.

8) Level gauge connections must not be used for taking samples.

8.2.2   H₂S Sampling and Analysis

This section is applicable to obtaining high pressure samples. Special considerations, such as extra sample(s) or purging, should be taken when obtaining low pressure samples (e.g., Boot, treater, stabilizer, acid gas).

Hydrogen sulphide (H_2S) is a reactive molecule, which presents challenges for sampling and analysis of gas mixtures containing it. Typically H_2S is lost during sampling (and analysis), resulting in underreporting of H_2S concentrations. Factors that affect representative sampling and analysis accuracy (i.e., the amount of H_2S lost) are

1)   presence of air, water, or other sulphur-containing molecules,

2)   presence of reactive or absorptive sampling container surfaces,

3)   presence of a liquid phase, which can absorb H_2S,

4) H_2S concentration,

5)   sample pressure and temperature,

6)   analysis method, and

7)   time lapse between sampling and analysis.

The amount of H_2S lost can be reduced by

- proper sample point selection, which minimizes the presence of contaminants such as air, water, and amines,

- using clean containers made of materials that minimize H_2S reactions or absorption, and

- minimizing the time between sampling and analysis.

Typical materials of construction for cylinders are stainless steel and aluminum. Inert coated cylinders, glass containers, and non-absorptive elastomer bags can be considered to further minimize H_2S degradation, especially for concentrations of H_2S less than 5000 ppm when moisture is present.

The choice of analytical technique also affects the amount of H_2S reported. Instrumental techniques, such as gas chromatography, are typically more precise than chemistry techniques, such as Tutweiler titrations or stain tubes. However, such instrumental techniques are often impractical for well site applications.

Therefore, consideration should be given to method limitations and sample degradation as they relate to the specific reporting requirements in determining the best approach.

See Table 8.1 for analysis technique comparison.

With the exception of ppm level concentrations of H_2S in the presence of moisture, a field H_2S determination and a laboratory GC analysis are recommended. These provide a degree of redundancy and a check of the field analysis. Above 5 per cent H_2S, the GC value is typically more reliable. Below 5 per cent H_2S, the higher of the two values should be used. Unexpectedly large variances between lab and field H_2S values need to be investigated.

Table 8.1. H₂S analysis technique comparison

Method	Lower detection limit	Advantages	Limitations
On-line GC	500 ppm	Real time, accuracy Minimal time lapse	Capital cost, ongoing maintenance
Lab GC	500 ppm	Precision, accuracy	Potential degradation during transport (varies with H₂S concentration)
Tutweiler (GPA C-1)	1500 ppm	On site	Titration apparatus, reagent quality, variability in operator technique, including visual endpoint detection, computations, mercaptan interference
Stain Tubes (GPA 2377)	1 ppm	On site	Poor precision (± 25%) Matrix effects, (see manufacturer's specifications)

Analysis by gas chromatography is the preferred method at higher H₂S concentrations.

For H₂S concentrations between 1500 and 5000 ppm, it is recommended that both stain tube and Tutweiler values be obtained if on-line GC is not used.

If high accuracy of low-level (below 1500 ppm) H₂S concentration is required, consideration should be given to using a low-level sulphur-specific detector, such as a GC sulphur chemiluminescence detector. The use of containers that minimize degradation and the time lapse between sampling and analysis is also recommended in these situations.

Refer to Appendix 5 for more detail on the analytical methods used in the industry for determining H₂S concentrations in gas samples.

8.2.3   Compositional Analysis of Natural Gas

The two recommended procedures for compositional analysis of natural gas are based on GPA Standard 2286-95: *Tentative Method of Extended Analysis for Natural Gas and Similar Gaseous Mixtures by Temperature Programmed Gas Chromatography* and GPA Standard 2261-00: *Analysis for Natural Gas and Similar Gaseous Mixtures by Gas Chromatography*.

If a thorough description of the C7+ fraction (molecular weight and density) is required, analytical methods based on GPA Standard 2286 are more accurate and preferred. Specifically, GPA Standard 2286 makes use of a high-resolution column and flame ionization detector to separate and quantify the heavier components (C7+), which is then used for calculation purposes. Extended analysis of natural gases is common but has not been fully standardized; therefore some inter-laboratory bias may occur.

If the C7+ properties are well defined or have been agreed upon by all affected parties, analytical methods based on GPA Standard 2261 are suitable. The principal advantage of the precut method specified in GPA Standard 2261 is that all of the C7+ components can be grouped together into a single sharp chromatograph peak. Grouping of the numerous heavy compounds results in more precise measurement of the combined peak area. The primary disadvantage of GPA Standard 2261 is the lack of information gained with respect to the composition of the C7+ fraction. Inherently, if the composition of the C7+ fraction is unknown, some agreed-upon physical properties must be applied for calculation purposes.

The GC C7+ calibration is also affected, which increases the uncertainty of the C7+ measurement and heating value computation.

If detailed information on C7+ physical properties is not available, default values can be applied, as in Table 8.2.

Table 8.2. Recommended default values for C7+ properties*

Component	Molecular mass (grams per mole)	Liquid density (kg/m³ at 15°C)	Heating value (MJ/m³)
C7+, Heptanes plus	95.00	735.0	195

* C7+ is a pseudo-compound. The values have in most cases had been found to adequately represent the heavier fraction of natural gas samples.

8.3 Gas Equivalent Factor Determination from Condensate

GEF is the volume of gas (10^3 m³ at standard conditions) that would result from converting 1.0 m³ of liquid into a gas.

GEF is used when there is a requirement to report the gas equivalent volume (GEV) of condensate and other hydrocarbon liquids to the ERCB.

The GEF of a liquid may be calculated by any one of three methods (see Appendix 6), depending upon the type of component analysis conducted on the liquid (by volume, mole, or mass fractions) and the known properties of the liquid.

Engineering Data

Specific constants are used in calculating the GEF. Absolute density of liquids should be used instead of relative density.

The examples in Appendix 6 present the different methodologies used to calculate the GEF. All physical properties are based on GPA Standard 2145-03 (2003 or later) published data.

1 kmol = 23.645 m³ @ 101.325 kPa and 15°C

8.3.1 Calculated Compositional Analyses

In some instances, representative sampling of a hydrocarbon stream is not possible or feasible because of economics, and calculation of a fluid composition is required, as described below:

Calculated Well Stream Analysis: It is not possible to accurately sample multiphase streams, so the composition of a recombined well stream must be determined by calculation. Such an analysis is typically not used for measurement, as it represents a multiphase fluid stream and most gas is measured as single phase. However, some companies use this analysis for calculation of gas volumes from wet- (multiphase) measured wells. Calculated well stream analyses are most commonly used in product allocation calculations.

Calculated Group Analysis: It is often difficult to accurately determine the average composition of fluids at a commingled group measurement point, as wells/sources to the group system flow at different rates and the composition is constantly changing. The recommended options for sampling these streams are on-line gas chromatographs or proportional sampling systems. However, if the recommended options are not practical or economical, a flow-weighted calculated analysis may be a viable option.

Calculated Single Analysis: Sometimes a single analysis cannot represent the composition for an entire measurement period. In such cases, multiple analyses of a single point must be combined to determine the composition for the period. An example of this is a sales gas stream where a proportional sample is taken weekly but a single composition for the month is required.

The principles to be followed for each of these calculated analyses follow.

8.3.1.1 Calculated Well Stream Compositional Analysis

This type of analysis applies to wells only and is meant to represent the hydrocarbon fluid composition produced from a well and/or delivered to a gathering system. In most cases, it represents the composition of hydrocarbons being produced from the reservoir. The calculation is a flow-weighted recombination of the hydrocarbon gas and liquid streams. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition. The gas and liquid flow rates used should be from the same day that the gas and liquid samples were obtained. Use of flow rates from a different period of time than the sample date can result in significant errors because both flow rates and composition will change with changes in process conditions (primarily temperature and pressure); applying flow rates from different periods does not recognize these changes.

Flow rates from the day of sampling should be used in determining recombined compositions, with the following exceptions:

- When the daily liquid-to-gas ratio is constant, volumes from an extended period (i.e., multiday, up to monthly) may be used.

- If some of the liquid stream is not recombined in a month (i.e., it dropped to tank), the composition (flow volume) of the liquids not recombined must be deducted from the initial recombined composition. This is typically done by recalculating the recombined composition with new flow rates, typically the flow rates for the month.

See the example in Appendix 7.

8.3.1.2 Calculated Group Compositional Analysis

This type of analysis is a flow-weighted representation of the hydrocarbon fluid composition produced from a group of wells or meter points. It is often used at commingled group points (inlets, compressors, certain process points) where it is difficult to obtain representative samples using spot sampling techniques. Ideally, proportional samplers should be employed in such situations. However, when proportional sampling is not practical or possible, a calculated group analysis can be determined based on the volume and composition of the wells/meters that flow to the commingled point. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition. The flow volumes used for each well/meter should be "actual" measured volumes for the period that the analysis is being calculated for, typically monthly.

For example, five gas wells producing from different pools with different composition deliver gas to a compressor station where the gas is measured. Accurate spot sampling at the compressor station is difficult due to changing flow rates at the wells. Using spot samples taken at the wells and monthly flow rates, the producer calculates a group analysis for the compressor station meter. Care must be taken when separator liquids are produced that all hydrocarbons are correctly accounted for, regardless of the phase.

See the example in Appendix 7.

8.3.1.3 Calculated Single Compositional Analysis

This type of analysis is a flow-weighted representation of the hydrocarbon fluid composition determined at a single sample point. It is typically used at sample points that have variable compositions and are sampled frequently (e.g., weekly) using spot or proportional sampling. Ideally, proportional samplers or gas chromatographs should be employed in such situations. However, when proportional sampling is not practical or possible, a calculated single analysis can be determined based on the volume and composition of a group of analyses at the sample point. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition. The flow volumes used for each sample should be "actual" measured volumes for the period that the analysis is representative of. For example, a producer takes spot samples of an inlet stream weekly because proportional sampling or on-line sampling is not practical. Using weekly flow rates, the producer calculates a monthly flow-weighted composition of the inlet stream.

See the example in Appendix 7.

8.4 Sampling and Analysis Frequency

Table 8.3 gives the analysis update frequency for gas and condensate streams. The sampling and analysis of condensate (if applicable) must be done at the same time as the gas sampling. The configurations shown in Figures 8.1 through 8.17 are examples. In each case, other similar configurations may also apply. Sampling frequencies are defined as follows:

Initial – *an analysis is required within the first six months of operation only, with no subsequent updates required.*

Monthly – *an analysis is required at least once per calendar month.*

Quarterly – *an analysis is required at least once per calendar quarter.*

Semiannually – *an analysis is required at least once every two calendar quarters.*

Annually – *an analysis is required at least once every four calendar quarters.*

Biannually – *an analysis is required at least once every eight calendar quarters.*

Calendar quarters *are January to March, April to June, July to September, and October to December.*

For example, for a biannual frequency, if the last sample was taken in July 2006, the operator has to take another sample by the end of September 2008 (end of the calendar quarter).

New gas and liquid samples must be taken for all new wells and measurement points by the end of the month following the first month of production. For the time period prior to receipt of a new composition, a substitute composition may be used for gas measurement and gas equivalent of liquid calculations. For wells, substitute compositions should be from a well producing from the same pool with similar separator operating conditions or from samples taken during well testing. Compositions taken during well tests should be carefully reviewed prior to use, as samples are typically taken at different conditions from those the well produces at and there are often contaminants in test samples (e.g., nitrogen, frac fluid). For non-well meters, the substitute composition should be as close to what is expected as reasonably possible. If the initial gas volume calculated by a substitute analysis is found to be in error by greater than 2% and the error volume is over $20 \times 10^3 \, \text{m}^3$/month, retroactive volumetric adjustments must be calculated using the initial gas composition.

Table 8.3. Sampling and analysis frequencies for various types of facilities

	Type of production battery/facility		Gas rate (10³ m³/d)	Sample and analysis type	Sampling point	Frequency
Gas wells/ batteries/ facilities	SE Alberta shallow gas zones/areas or coalbed methane well with minimal water (see 8.4.1)		N/A	Gas only	Group meter	Biannually
	Gas proration outside SE Alberta shallow gas zones /areas or effluent proration battery (see 8.4.2 & 8.4.3)		N/A	Gas/condensate	Test meters	At time of testing
					Group meter	Annually
	Multiwell group battery or single-well battery (including coalbed methane well/battery) with no condensate or oil (see 8.4.4)		N/A	Gas only	Per meter	Annual first year, then biannually
	Multiwell group battery or single-well battery with condensate or oil (see 8.4.5)		> 16.9	Gas/condensate	Per meter	Annually
			≤ 16.9	Gas/condensate	Per meter	Biannually
	Gas storage schemes, injection and withdrawal phase (see 8.4.6)			Gas	Per injection/ production meter	First month, then semi-annually
	Gas cycling schemes (see 8.4.7)	Injection		Gas/condensate	Per injection meter	Per approval or source requirement (if not in approval)
		Production			Per production meter	Per approval or semi-annually (if not in approval)
	Gas sales/delivery (see 8.4.8)			Gas only	Per meter	Annually
	Gas plants (see 8.4.9)			Gas/condensate	Per accounting meter	Semiannually
	Gas gathering systems (see 8.4.9)		> 16.9	Gas/condensate	Per inlet meter	Annually
			≤ 16.9	Gas/condensate	Per inlet meter	Biannually
Conventional oil wells/ batteries	Single-well/ multiwell group battery (see 8.4.10)	Flared		Gas only	Per meter	Initial
		Conserved	> 16.9	Gas only	Per meter	Annually
			≤ 16.9	Gas only	Per meter	Biannually
	Multiwell proration battery	Primary production and water flood (see 8.4.11)	> 16.9	Gas only	Per test/group meter per pool	Annually
			≤ 16.9	Gas only	Per test/group meter per pool	Biannually
		Miscible/immiscible flood (see 8.4.12)	Production	Gas only	Per test/group meter	Per approval or quarterly (if not in approval)
			Injection		Per meter	Per approval or monthly (if not in approval)
Heavy oil / crude bitumen batteries (see 8.4.13)	Single-well/ multiwell group battery		≤ 2.0	Gas only	Per well	Initial
			> 2.0	Gas only	Per meter	Biannually
	Multiwell proration battery		≤ 2.0	Gas only	Per meter/ Per pool	Initial
			> 2.0	Gas only	Per test/group meter	Biannually

Licensees must ensure that analysis data are used to update volumetric calculations by the end of the month following the receipt of the analysis report. The only exception is for effluent wells, for which the analysis must be updated by the end of the second month following receipt of the analysis report. If sampling and analysis are conducted for other purposes (e.g., joint venture and allocation agreements) more frequently than required by this directive, the licensee must use those data to update volumetric calculations.

8.4.1 Southeastern Alberta Shallow (SEA) Gas Zones/Areas or Coalbed Methane Wells (with Minimal Water)

In the configuration of Figure 8.1 of shallow gas wells or coalbed methane (CBM) wells with water production not more than $0.01 \, m^3 / 10^3 \, m^3$ gas, analyses must be updated biannually at each group meter (Meter A). No sampling and analysis are required at the test meter or well. Density and component analysis data from the group meter sample point may be used for test meter calculations.
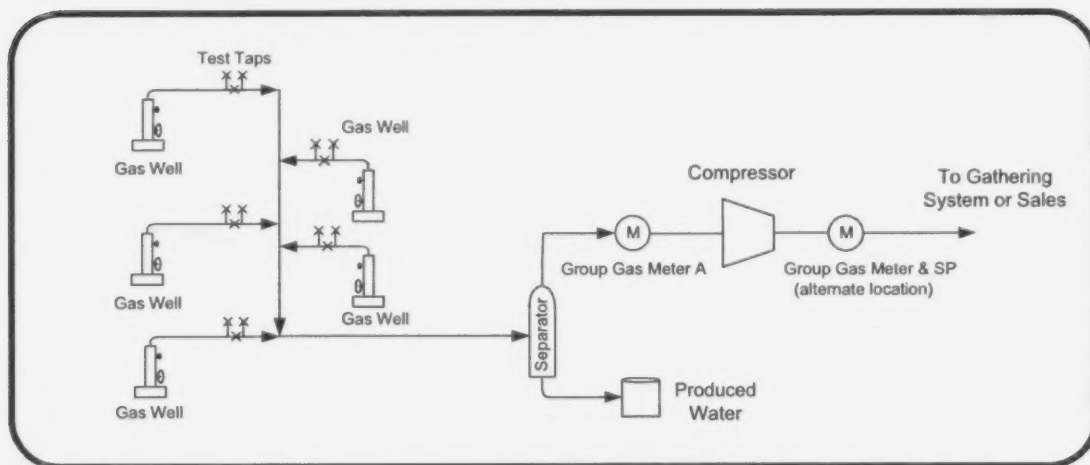
Figure 8.1

For coalbed methane wells that have been fractured or stimulated using a gaseous medium, gas samples must be taken monthly until the composition stabilizes and then biannually.

If these zones are commingled with gas from other outside zones, sampling must be done on a per pool/zone basis for CBM or per commingled SEA pool(s) basis from a representative well within the pool(s) biannually.

For exceptions on sampling for these pools/zones, see Section 8.4.2.

8.4.2 Gas Proration Outside SE Alberta Shallow Gas Zones/Areas (Excluding Effluent Proration Battery)

In the configuration in Figure 8.2, gas density and composition must be updated annually at each group meter (Meter A) and at each well during a test. If condensate at the group measurement point is recombined and delivered to a gas plant, the condensate GEF must be updated annually.
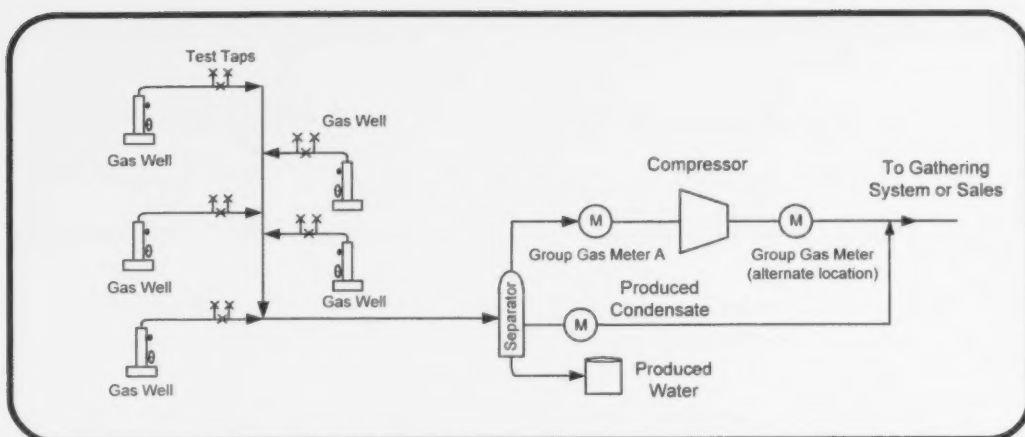
Figure 8.2

Exceptions

1) A licensee is not required to update the analyses where three consecutive gas relative density (RD) determinations conducted at the specified determination frequency or, alternatively, no more frequently than once per year are all within ±1.0 per cent of the average of the three RDs (see Example 8.1). In this situation, there is no need for an application to be submitted to the ERCB. Records and data in support of this exception must be retained by the licensee and made available to the ERCB upon request. Notwithstanding this exception, the licensee must update the gas analyses when changes are made to producing conditions that could affect the gas density by more than ±1.0 per cent of the average of the three qualifying RDs.

2) The gas analyses determined at the group measurement points may be used for the test meters, provided that all wells are from the same pool.

Example 8.1

		Meter A
Sample date	RD	Difference from average
June 03, 2000	0.583	- 0.29%
June 09, 2001	0.586	+ 0.22%
June 06, 2002	0.585	+ 0.05%
Average	0.5847	

In this example, Meter A would be exempt from the requirement for future updates as the three consecutive RDs are within ±1.0 per cent of the average of the three RDs.

8.4.3   Multiwell Effluent Proration Battery

Wellhead Testing Sampling Points

In the configuration in Figure 8.4, gas analyses, condensate composition, and GEF must be updated at the time of testing each effluent well and annually at the group gas and condensate meters (see Figure 8.3). The gas analysis to be used for volumetric calculation at the effluent meter is as follows:

- Option 1: Use the separated gas analysis from the ECF-WGR test; or

- Option 2: Use the recombination of the gas analysis and the condensate analysis from the ECF-WGR test.

Note that all wells within the effluent battery must use the same analysis option.

Exceptions

A licensee is not required to update analyses at the group gas and condensate metering points if

- three consecutive gas RD determinations conducted at the specified determination frequency or no more frequently than once per year are all within ±1.0 per cent of the average of the three RDs (see Example 8.1), and

- the daily average liquid condensate volume is less than or equal to 2.0 m^3/d for all reporting months for the previous three years and/or the GEV of the condensate is less than or equal to 2.0 per cent of the recombined total monthly gas volume.

In these situations, there is no need for an application to be submitted to the ERCB. Records and data in support of these exceptions must be retained by the licensee and made available to the ERCB upon request. Notwithstanding these exceptions, the licensee must update the gas analyses when changes are made to producing conditions that could affect the gas RD by more than ±1.0 per cent of the average of the three qualifying RDs, and the licensee must update the condensate analyses if the liquid condensate volume or GEV percentage increases beyond the qualifying limits.

Figure 8.3. Typical multiwell effluent proration battery
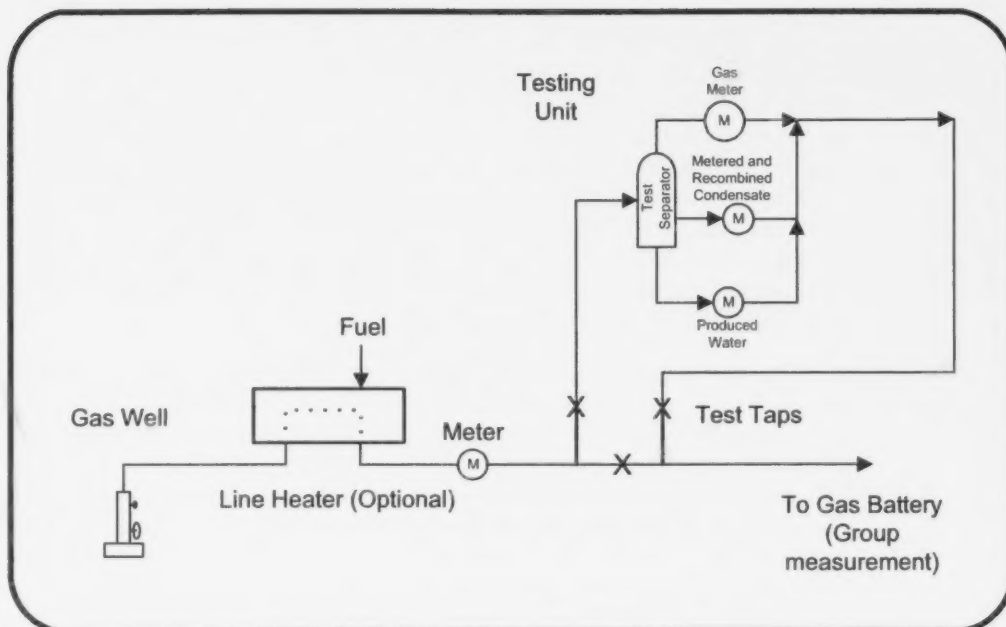
Figure 8.4. Typical effluent testing configuration

8.4.4 Single Gas Well or Multiwell Group Gas Battery and Coalbed Methane Well with No Condensate or Oil

A "multiwell group" is a number of measured single gas wells all connected to a common delivery point.

For the configuration in Figure 8.5, the gas analysis must be updated within the first year and then biannually at each well meter (Meter A). For coalbed methane wells that have been fractured or stimulated using a gaseous medium, gas samples must be taken monthly until the composition stabilizes and then biannually.
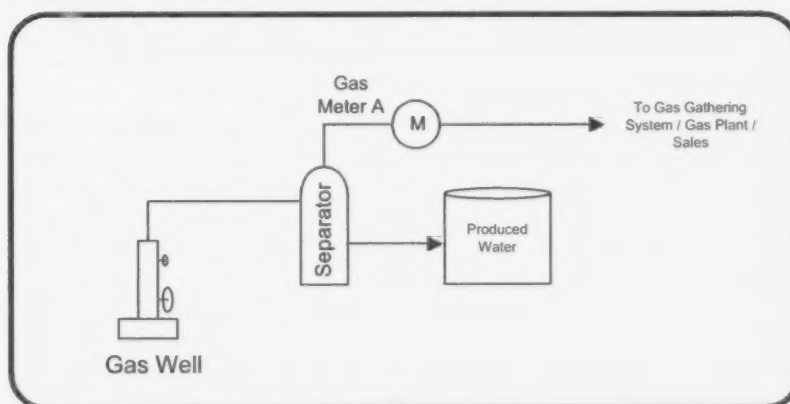
Figure 8.5

Exceptions

1) A licensee is not required to update the analyses if three consecutive gas analyses conducted at the specified determination frequency or, alternatively, no more frequently than once per year are all within ±1.0 per cent of the average RD of the three analyses (see Example 8.1). In this situation, there is no need for an application to be submitted to the ERCB. Records and data in support of this exception must be retained by the licensee and made available to the ERCB upon request. Notwithstanding this exception, the licensee must update the gas analyses when changes are made to producing conditions that could affect the gas analysis by more than ±1.0 per cent of the average RD of the three qualifying analysis.

2) A representative analysis for all wells producing to a common gathering system or facility from a common pool can be used if the RD of all common-pool wells are within 2.0 per cent of the average analysis of those wells. Gas analyses must initially be obtained for all the common-pool wells to arrive at the average analysis. Subsequent analyses can be made on 25 per cent or at least four wells from the pool (whichever is greater) at the frequency stated in this directive, provided that the RD variance remains within the 2.0 per cent limit of these wells (see Example 8.2). Should the variance exceed this limit, this exception is revoked and biannual analyses must be determined for each measurement point.

Example 8.2

Consider an 8-well pool producing gas under the configuration below:

Well	RD	Difference from average
11-14	0.602	-0.99%
10-16	0.610	+0.33%
10-21	0.602	-0.99%
9-27	0.616	+1.32%
11-30	0.608	0.00%
6-31	0.616	+1.32%
11-32	0.606	-0.33%
11-16	0.604	-0.66%
Average	0.608	

In this case, it is acceptable to use the analyses from the well with the RD closest to the average (11-30) for all well meters, as all RDs are within ±2.0 per cent of the average of all well RDs. The analysis must then be updated biannually for at least four wells from the pool. This exception will remain in place, provided that all four well RDs continue to be within ±2.0 per cent of the average of all the updated RDs. When this criterion is not met, analyses must revert to biannual updates for all wells.

Permanent exception on a pool basis would be available where the updated average RD meets the criterion of Exception 1. Where practical, the ERCB expects the same wells to be used to arrive at the average RD used in pursuit of this exception.

8.4.5 Single Gas Well or Multiwell Group Gas Battery with Condensate or Oil

A "multiwell group" is a number of measured single gas wells all connected to a common delivery point.

For gas wells producing condensate in Figure 8.6, the frequency of sampling and analysis for gas and condensate depends upon the gas flow rate through Meter A plus the GEV of condensate. If the flow rate exceeds $16.9 \times 10^3 \, m^3/d$, the frequency is annual. If the flow rate is less than or equal to $16.9 \times 10^3 \, m^3/d$, the frequency is biannual. The flow rate is to be based on a monthly average.
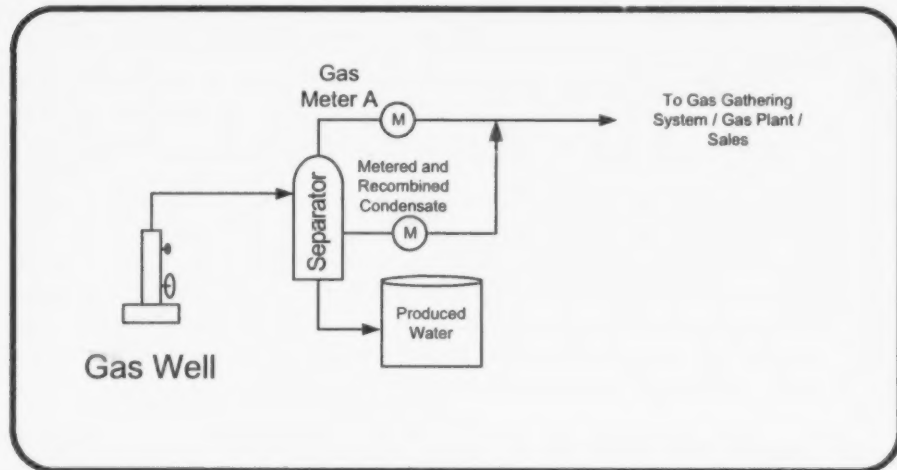
Figure 8.6. Condensate production

For gas wells producing oil in Figure 8.7, the sampling and analysis of oil/emulsion streams to determine relative oil and water content must conform to the requirements in Section 6.4.3. The gas sampling frequency is the same as for a gas well producing condensate, except that the total gas flow rate does not include GEV of oil/emulsion.
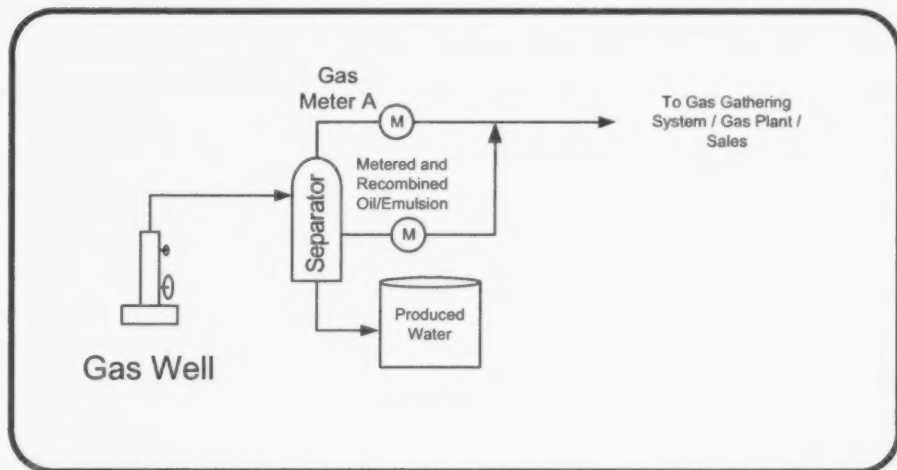
Figure 8.7. Oil production

Exceptions

1) For gas wells producing condensate, a licensee is not required to update the gas and condensate (if applicable) analyses at the metering points if

   • three consecutive gas RD determinations conducted at the specified determination frequency or no more frequently than once per year are all within ±1.0 per cent of the average of the three RDs (see Example 8.1), or

   • daily average liquid condensate volume is less than or equal to 2.0 m^3/d for all reporting months for the previous three years and/or the GEV of the condensate is less than or equal to 2.0 per cent of the recombined total monthly gas volume.

   In these situations, there is no need for an application to be submitted to the ERCB. Records and data in support of these exceptions must be retained by the licensee and made available to the ERCB upon request. Notwithstanding these exceptions, the licensee must update the gas analyses when changes are made to producing conditions that could affect the gas RD by more than ±1.0 per cent of the average of the three qualifying RDs, and the licensee must update the condensate analyses if the liquid condensate volume or GEV percentage increases beyond the qualifying limits.

2) A **representative** analysis for all wells producing to a common gathering system or facility **from a common pool** may be used if the RDs of all common-pool wells are within 2 per cent of the average RD of those wells. Gas analyses must initially be obtained for all the common-pool wells to arrive at the average RD. Subsequent analyses may be made on 25 per cent or at least four wells from the pool (whichever is greater) at the frequency stated in this directive, provided that the RD variance remains within the 2 per cent limit of these wells (see Example 8.2). Should the variance exceed this limit, this exception is revoked and biannual analyses must be determined for each measurement point.

8.4.6 Gas Storage Schemes

For the configuration in Figure 8.8, there are two phases to consider:

Injection Phase

If the injection gas only comes from a single source, an annual common stream sample analysis may be used for all injection meters, and no individual well injection analyses are required.

If there are multiple injection gas sources, sample analysis is required at each source stream and at each well injection measurement point. In this case, the minimum analysis frequency for injection meters are semiannual, however, a continuous proportional sampler or a gas chromatograph should be installed to provide more accurate compositions for gas volume calculations.

Withdrawal Phase

During each withdrawal phase, analyses must be updated at each well's (A, B, and C) production meters within the first month and semiannually thereafter if necessary.
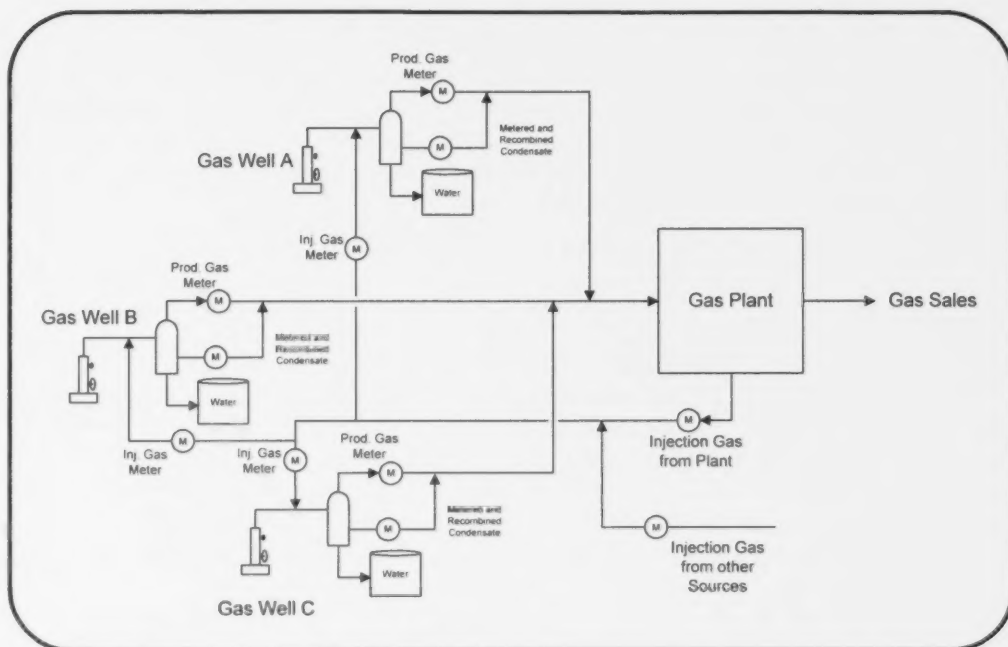
Figure 8.8

8.4.7 Gas Cycling Scheme

In the configuration in Figure 8.9, analyses must be updated at each well meter, A, B, and C, and the injection well meter in accordance with the specific scheme approval. If there are no frequencies specified in the approval, the well meters must have analyses updated semi-annually and the gas injection meter(s) must have analyses updated in accordance with the source requirements (e.g., semiannually for gas plant gas).
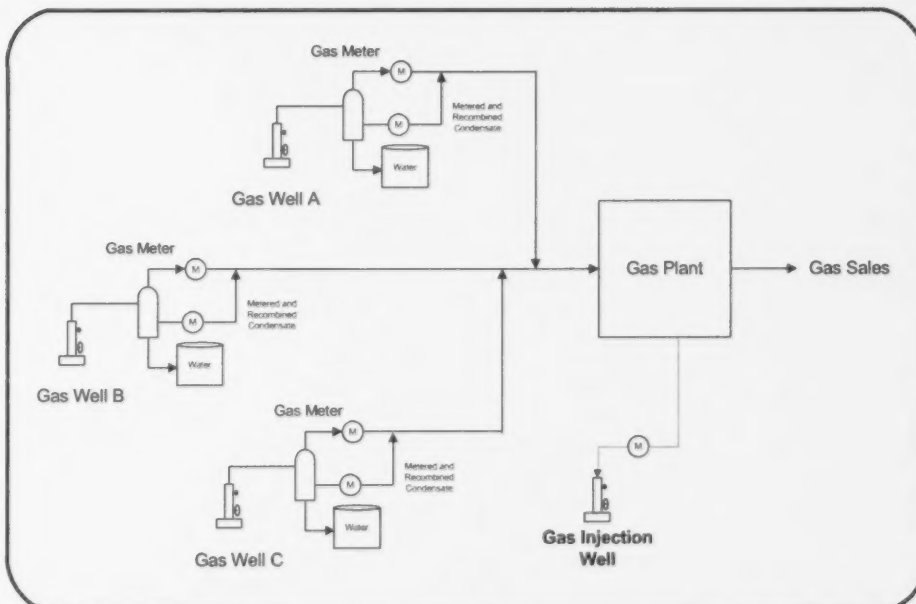
Figure 8.9

8.4.8 Gas Sales/Delivery

In the configuration in Figure 8.10, gas sales/delivery in this context will typically be clean, processed sales gas that is delivered out of a gas plant or a facility into a transmission pipeline. The measurement at this point determines the gas volumes upon which royalties will be based. In some cases, this type of gas may be delivered to other plants for further processing or fuel or to injection facilities.
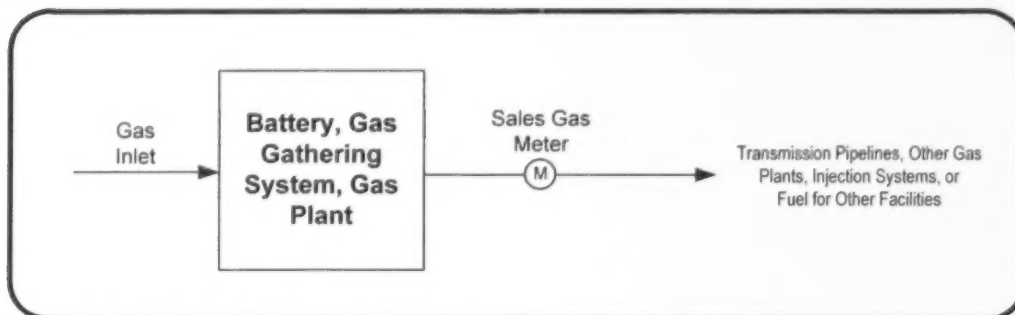
Figure 8.10

If a meter is used to determine the sales gas/delivery point volume from a battery, gas gathering system, or gas plant, the minimum gas analysis frequency is annual. However, a continuous proportional sampler or a gas chromatograph should be installed to provide more accurate analyses for the gas volume calculation.

8.4.9 Gas Plants and Gas Gathering Systems

In the configuration in Figure 8.11, only one sample point is required for common gas streams, such as sales gas, which may also be used for fuel, injection, and sales gas flare. Inlet gas sample may be used for inlet gas flare.

The frequency for sampling and analysis is as follows unless a different frequency has been specified in site-specific approvals, such as gas cycling or miscible/immiscible flood schemes, or for heavy oil/crude bitumen gas production. For gas sales measurement point sampling frequency, see Section 8.4.8.

Gas Plant

The minimum frequency for updating analyses at all accounting meters within a gas plant is semiannual. Inlet condensate is reported as a GEV, so analyses are required. High-vapour pressure liquids, such as pentanes plus and NGL, are to be reported as liquid volumes on the PRA, which will then perform the GEV calculation automatically using standard factors for plant balancing.

Gas Gathering System

The minimum frequency for updating analyses at all accounting meters within a gas gathering system (GGS) is annual for all flow rates that exceed $16.9 \ 10^3 \ m^3/d$. If the flow rate is less than or equal to $16.9 \ 10^3 \ m^3/d$, the frequency is biannual. The flow rate is to be based on a monthly average. Condensate volumes recombined with gas for delivery to other facilities must be reported as GEV, so analyses are required for updating GEFs. Where condensate is delivered out of a GGS without further processing, it is reported as a liquid volume, but analyses for GEV calculation purposes are required for reporting to the PRA.
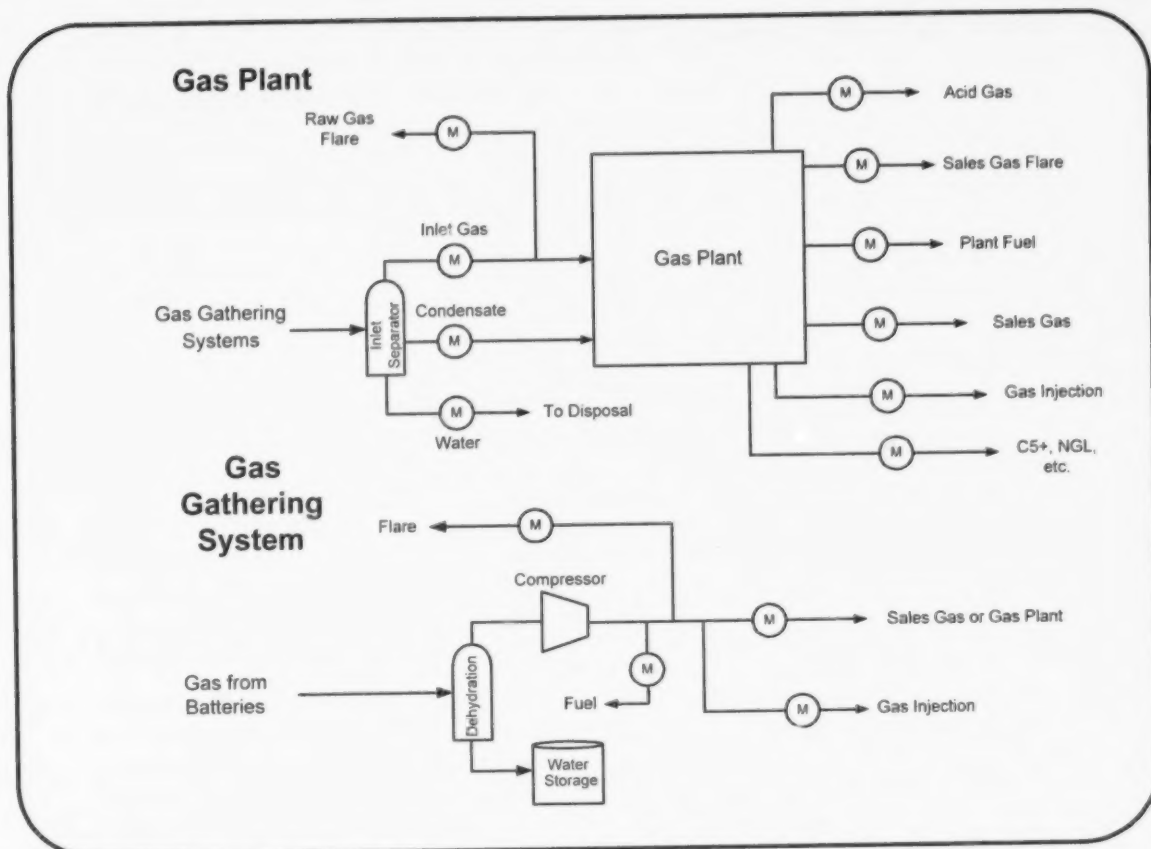
Figure 8.11

8.4.10 Conventional Oil Facilities

In the configuration in Figure 8.12, if all solution gas (net of lease fuel) is flared, an initial representative gas analysis is required. If gas is conserved, gas analysis updates are required. If the average flow rate exceeds 16.9 10^3 m^3/d, the frequency is annual. If the average flow rate is less than or equal to 16.9 10^3 m^3/d, the frequency is biannual.
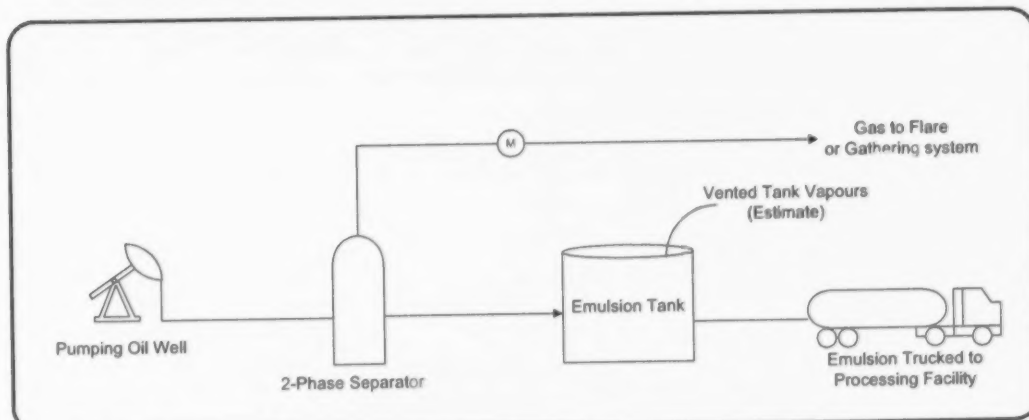
Figure 8.12. Single-well or multiwell group oil battery

8.4.11 Multiwell Proration Oil Battery

In the configuration in Figure 8.13, the gas analyses must be updated at the test meters (A and B) biannually for maximum test gas rates up to 16.9×10^3 m³/d or annually if the maximum test gas rates exceed 16.9×10^3 m³/d.
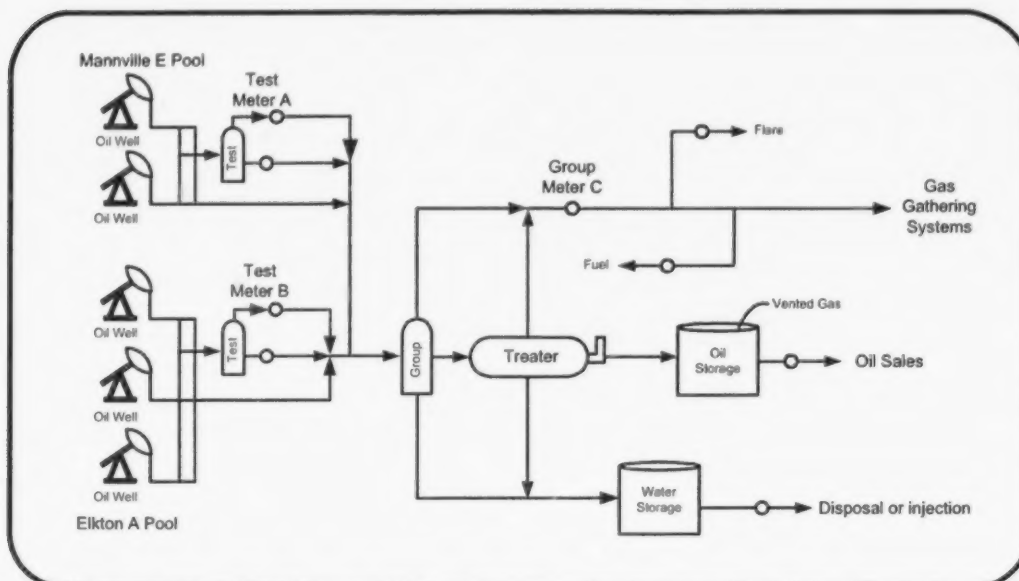
Figure 8.13. Primary production/water flood

It is acceptable to use the gas analysis from a single representative well for all wells within a single pool. If wells from more than one pool are directed through the same test separator, an analysis must be obtained for each pool.

The gas analysis at meter C must be updated annually for gas flow rates exceeding 16.9×10^3 m³/d or biannually if the total rate through the meter is less than or equal to 16.9×10^3 m³/d (based on the monthly average flow rate).

Example 8.3

Consider a 5-well proration battery with two wells producing from the Mannville E Pool and three wells producing from the Elkton A Pool. Battery gas production is gathered and conserved.

Pool	Well	Satellite meter	Test gas rate (10^3 m³/d)
Mannville E	10-14	Meter A	4.2
Mannville E	10-16	Meter A	6.8
Elkton A	10-21	Meter B	18.0
Elkton A	9-27	Meter B	12.0
Elkton A	10-30	Meter B	6.5
		Total rate for Meter C =	47.5

A gas analysis must be established for the Mannville E Pool, as a minimum using either the 10-14 or 10-16 well, and updated biannually at meter A, as the maximum rate through meter A for the Mannville E pool wells is less than 16.9 10^3 m^3/d.

A gas analysis must be determined for the Elkton A Pool at meter B, as a minimum using any one of the three wells, and updated annually, as the maximum rate through meter B for the Elkton A pool wells is greater than 16.9 10^3 m^3/d.

The gas analysis at meter C must be updated annually, as the flow rate through the meter exceeds 16.9 10^3 m^3/d.

Exception

If the total battery gas, net of lease fuel, is flared, an initial pool gas analysis must be determined at meters A and B. Updates of the gas analysis at meter C, at the annual or biannual frequency as determined by the gas flow rate through the meter, is only required if the gas directed through meter C originates from multiple pools. If the gas directed through meter C originates from a single pool, no updates are required subsequent to the initial analysis. However, this exception is revoked as soon as the gas is conserved, and gas analyses must be performed according to the frequencies specified above.

8.4.12 Miscible/Immiscible Flood

In the configuration in Figure 8.14, analyses must be updated at each test and group meter and the injection well meter in accordance with the specific scheme approval. If there are no frequencies specified in the approval, the sampling frequency must be updated in accordance with the requirements in *IL 92-05*. That is, the test and group meters must have analyses updated quarterly and the injection meter(s) must have analyses updated monthly.
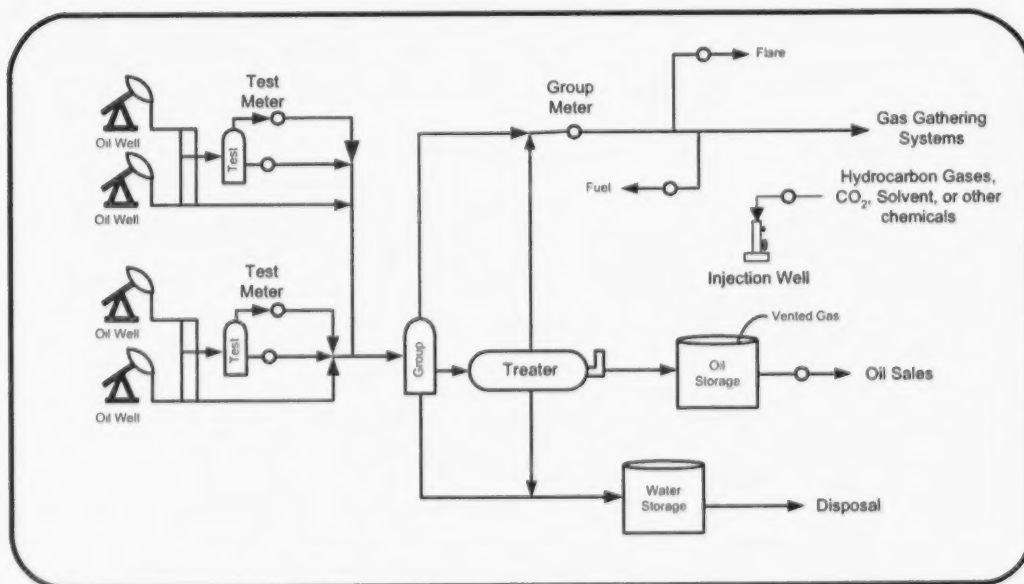
Figure 8.14

8.4.13   Heavy Oil/Crude Bitumen Facilities

Heavy oil/crude bitumen production at a single-well (Figure 8.15) or multiwell group battery (Figure 8.16) typically involves directing all production to a tank without using a separator or gas meter. In such cases, gas production may be estimated using a GOR. If a meter is used to measure gas for the purposes of conducting GOR tests or continuous gas production measurement, an initial gas analysis is required. An analysis from a comparable well producing from the same pool may be used if a meter will be used to measure gas to determine GOR.
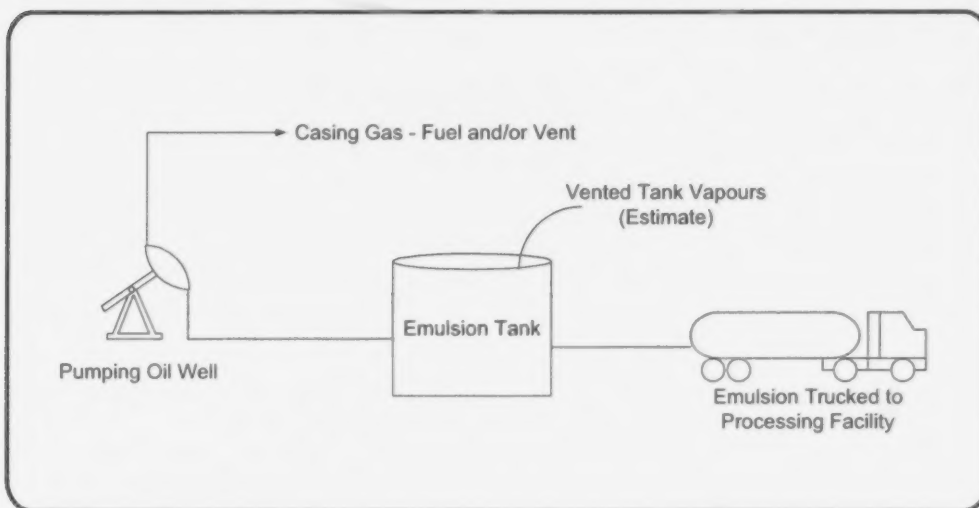
Figure 8.15. Heavy oil/crude bitumen single-well battery
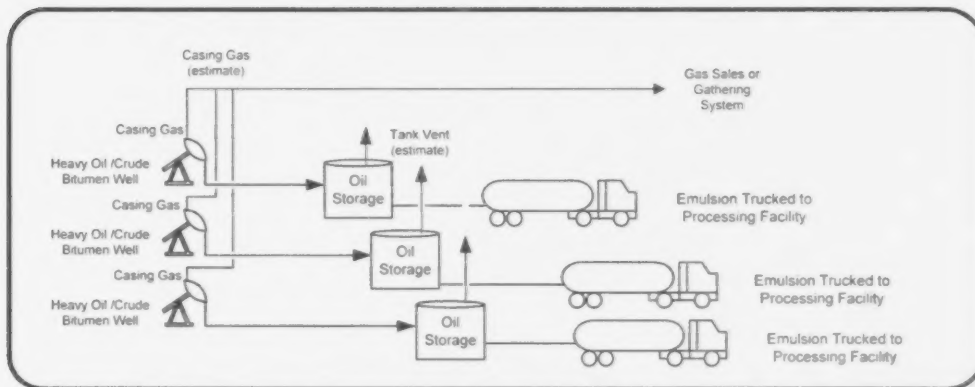
Figure 8.16. Heavy oil/crude bitumen multiwell group battery

If a GOR is determined by methods other than using gas measurement, an initial gas analysis is not required.

If a meter is used to measure gas on a continuous basis, biannual analysis updates are required.

Heavy oil/crude bitumen production at multiwell proration batteries (Figure 8.17) may involve directing all production to tanks without using separators or gas meters, but if

combined gas volumes meet the economic requirements in *Directive 060*, the gas must be gathered and conserved.
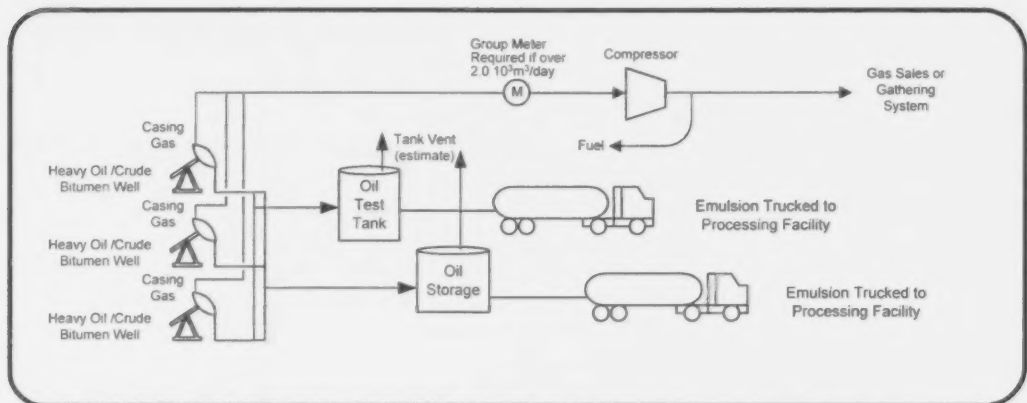
Figure 8.17. Heavy oil/crude bitumen multiwell proration battery

If a meter is used to measure gas for the purposes of conducting GOR tests, an initial gas analysis is required. If a GOR is determined by methods other than using gas measurement, an initial gas analysis is not required.

If a meter is used to measure gas on a continuous basis, biannual analysis updates are required.

9 Cross-Border Measurement

This section presents the measurement requirements for all upstream and midstream oil and gas production crossing a provincial or territorial border.

9.1 General Requirements

For those facilities receiving non-Alberta production and/or delivering Alberta production to another jurisdiction either by trucking or pipeline, including pipelines under the National Energy Board (NEB) jurisdiction, each jurisdictional production stream must be isolated and measured prior to commingling. The delivery point measurement standards for Alberta and that of the other jurisdictional authority(ies) must be followed, unless site-specific approval from the ERCB and the other jurisdictional authority(ies) has been obtained. All streams must be isolated and measured or estimated according to requirements in this directive. This could include production and gathering systems, all fuel, flare, and vent. If the measurement or other equipment requirement for delivery point measurement of hydrocarbon and related fluids from any jurisdiction is different from the Alberta requirements, the higher requirements, such as frequency and accuracy, between the jurisdictions must be followed.

"Non-Royalty Exempt Fuel Gas" usage at a cross-border oil and gas processing facility must be separately determined and measured if it is over $0.5 \ 10^3 \ m^3/d$ for each jurisdiction. If the usage is for production from both jurisdictions or site-specific approval is obtained from both jurisdictions involved, no separate fuel gas metering is required. For example, a compressor used only for gas coming from another jurisdiction into Alberta must be metered separately at the cross-border facility and the fuel gas use for other equipment processing commingled production or the entire facility must be measured with another meter.

9.2 Cross-Border Sampling Requirements

Except where otherwise noted, the gas and liquid sampling equipment and methodology must follow the requirements set out in Section 8: Gas and Liquid Sampling and Analysis.

Spot or grab samples are acceptable for obtaining gas and liquid analyses, provided the uncertainty requirements in Section 1 of *Directive 017* are fulfilled.

When the uncertainty requirements cannot be met, consider

- more frequent spot sampling for calculated analysis,
- the use of proportional samplers, or
- the use of gas chromatographs or other continuous analyzers.

9.3 Cross-Border Measurement Points

Figures 9.1 through 9.9 are some (not an exhaustive set) of the production scenarios to determine if a specific circumstance is considered "cross border." Each scenario applies as well if the flow is in the opposite direction. There must be only one cross-border measurement point for each pipeline crossing the provincial boundary unless site-specific approval is obtained from both jurisdictions involved.

The cross-border measurement point can be on either side of the jurisdictional border before commingling with any fluids from another jurisdiction. Measurement-by-difference rules apply in all cases where there is measured production going into a proration battery.
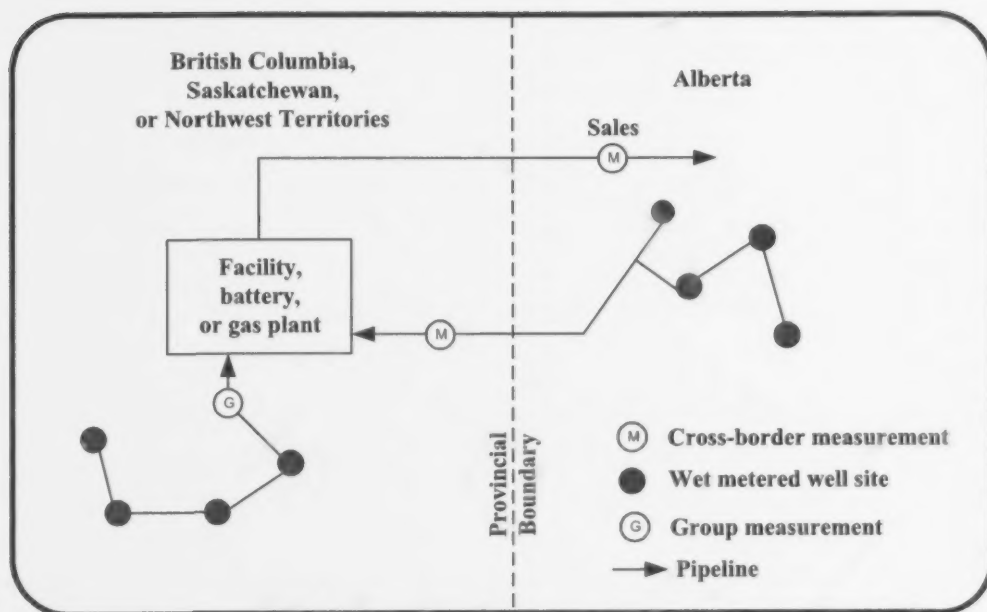
Figure 9.1. Wet gas measured wells to an out-of-province location (non-common pool)

For cross-border common pools producing from one or more jurisdictions, if the surface facility is located in one jurisdiction and the well production as defined by the bottomhole location is in another jurisdiction, delivery point measurement of the production is required (see Figure 9.2). The production from this well must be reported as delivered to the other jurisdiction where the surface facility is located.
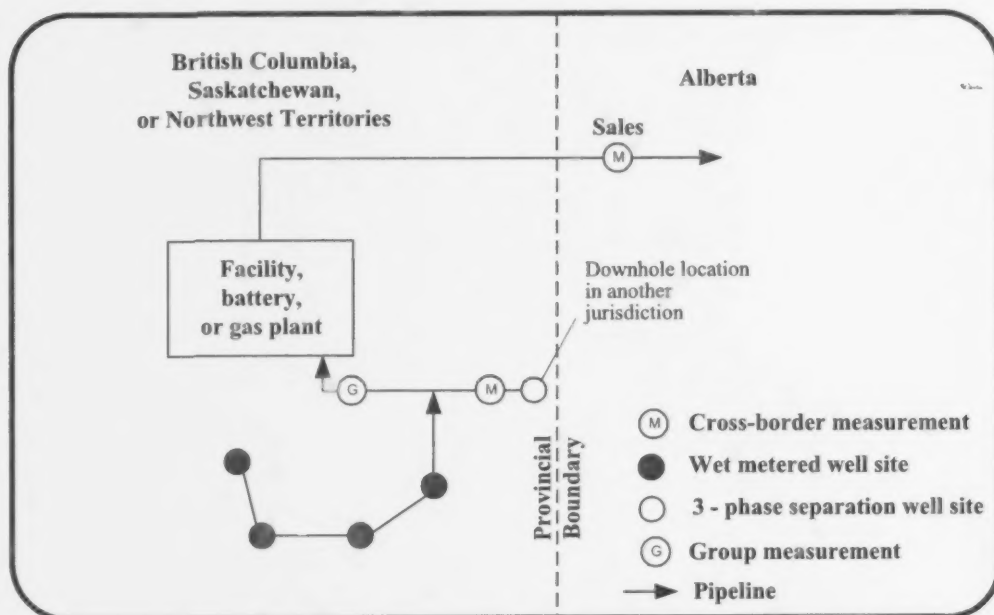
Figure 9.2. Gas gathering system of wet gas measured well with measured well downhole location in another jurisdiction
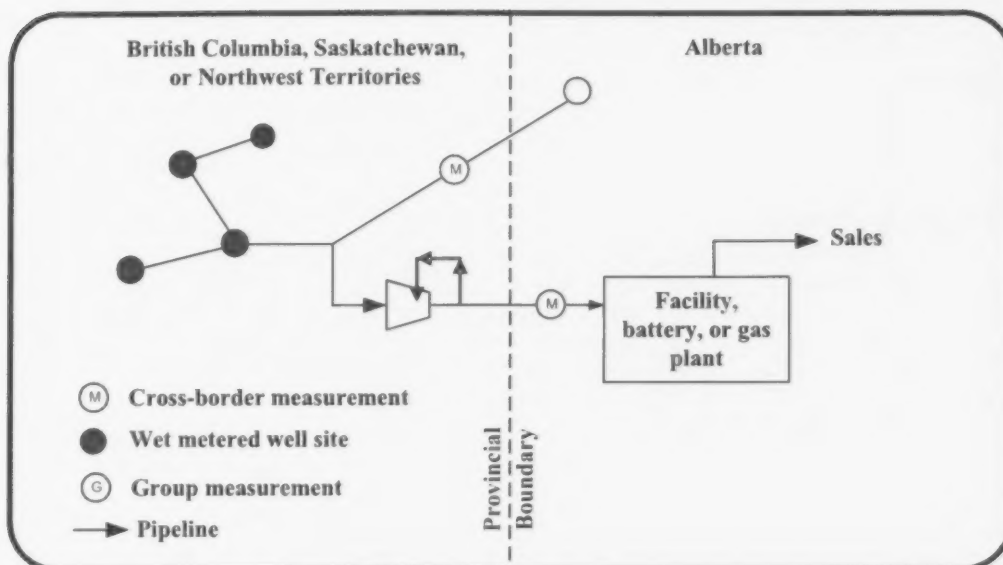
Figure 9.3. Multiple jurisdictional crossing
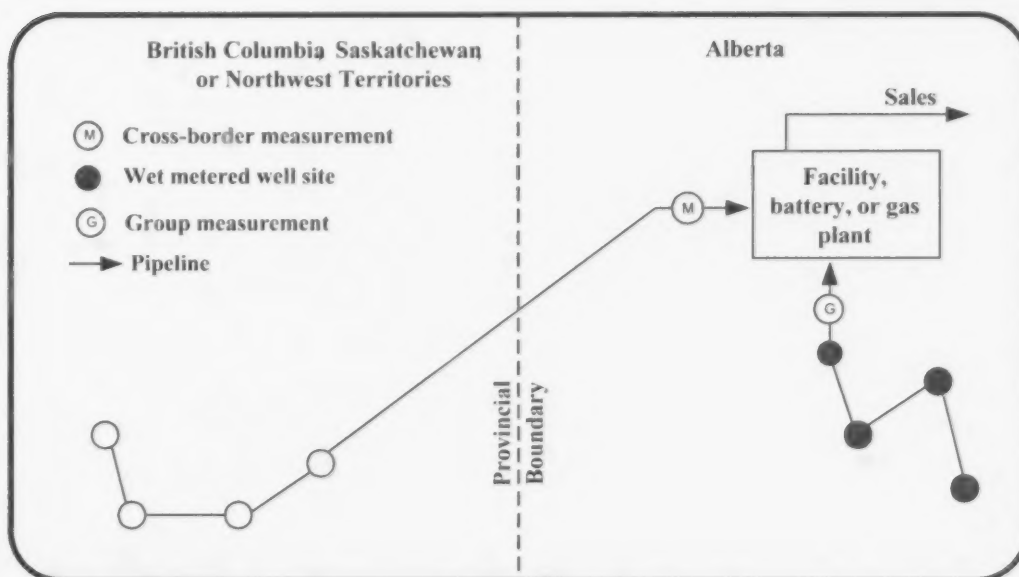
Figure 9.4. Measured gas source from an out-of-province location
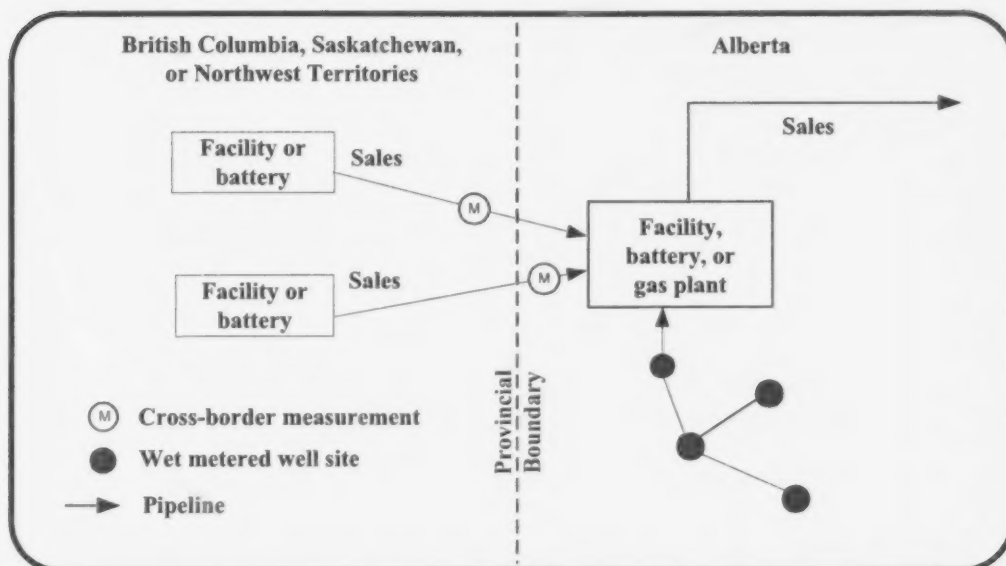
Figure 9.5. Sales gas source from an out-of-province location
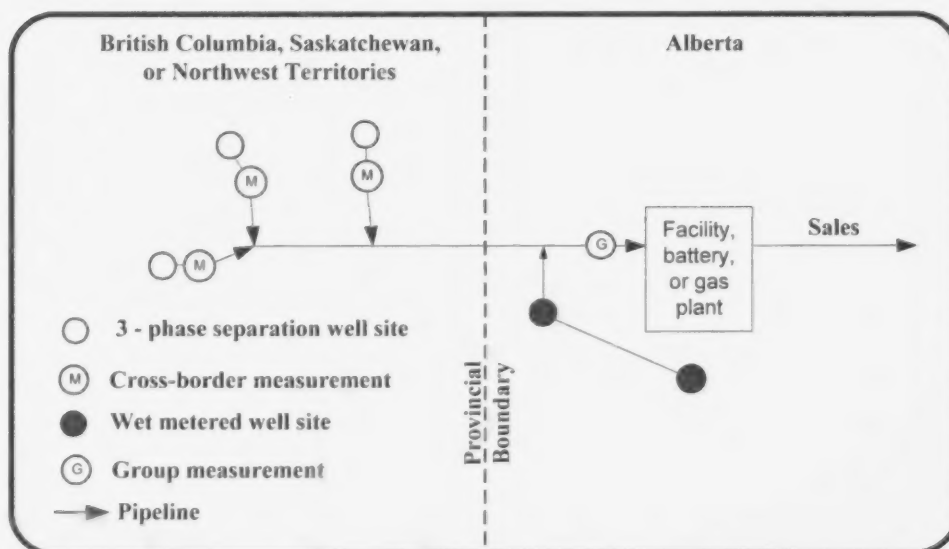
Figure 9.6. Measured sources from an out-of-province location

For Figure 9.6, the three-phase separation at the well site can be designed to delivery point measurement requirements without another cross-border measurement point.

Measurement-by-difference rules applies in Figures 9.1, 9.2, 9.4, 9.5, and 9.6 unless the wet metered wells have a group measurement point prior to commingling with the measured gas source(s).
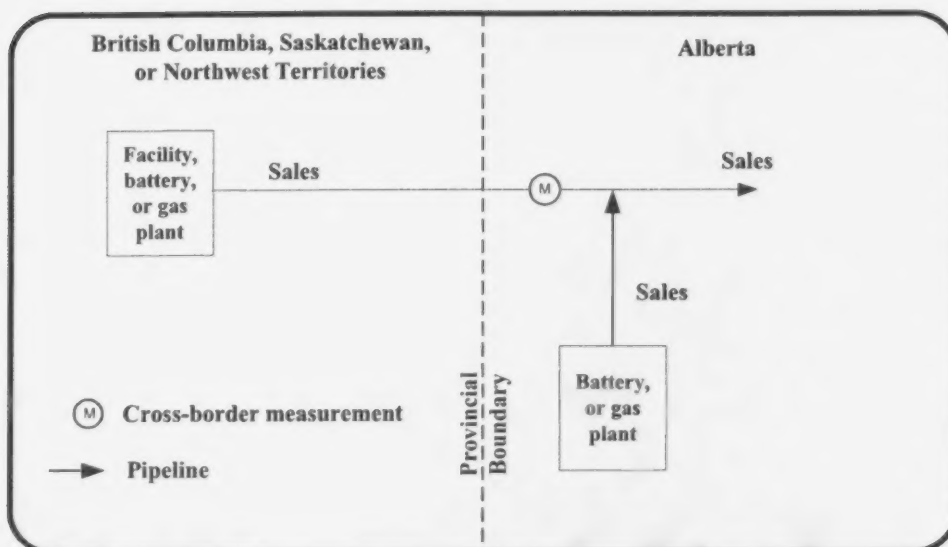
Figure 9. 7. Sales oil or gas source from an out-of-province location
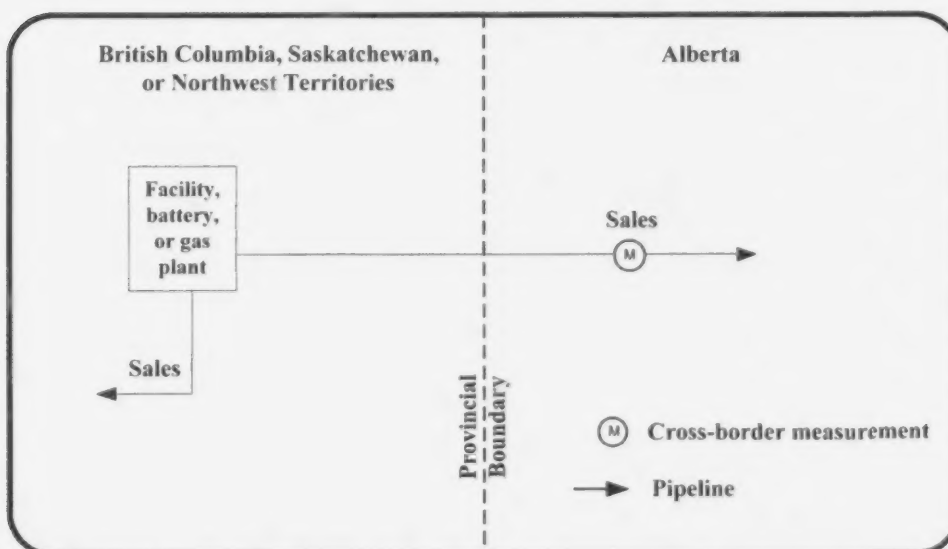
Figure 9.8. Sales oil or gas source from an out-of-province location
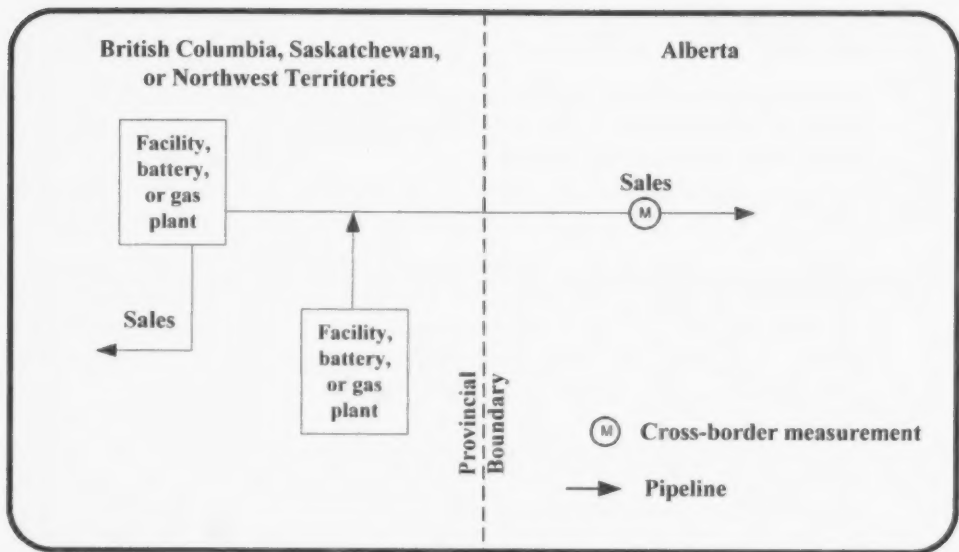
Figure 9.9. Sales oil or gas source from an out-of-province location

10 Trucked Liquid Measurement

This section presents the requirements for trucked liquid measurement from oil and gas production facilities to another facility or sales. Applicable liquids include oil, condensate, water, NGLs, propane, and butane.

10.1 General Requirements

Crude oil or condensate may be found in association with water in an emulsion. In such cases, the total liquid volume of the trucked load must be measured, and the relative volumes of oil and water in the emulsion must be determined by obtaining and analyzing a representative sample of the emulsion using a product analyzer such as a water-cut monitor or a Coriolis meter's density measurement (where applicable).

A licensee must accurately measure produced liquids/emulsion volumes by using tank gauging, a weigh scale, or a meter, unless otherwise stated in this directive. The delivery point measurement requirements must be met for all trucked liquids unless the exception conditions in this section are met or site-specific approval from the ERCB has been obtained.

The ERCB will consider a truck liquid measurement system to be in compliance if the following base requirements are met. The ERCB may stipulate additional or alternative requirements for any specific situation, based on a site-specific assessment.

All delivery point meters must be proved in accordance with Section 2. LACT meters may use the proving procedure in API-MPMS, Chapter 4: Proving Systems, instead of the Section 2 procedure.

10.1.1 Reporting Requirements

Monthly oil, condensate, and water volumes for well(s) (production) and battery(s) (production, receipts, and dispositions) must be reported as the number of cubic metres rounded to the nearest tenth of a cubic metre (0.1 m^3). Measured volumes must be corrected to 15°C and at the greater of 0 kPag or equilibrium vapour pressure at 15°C. See Section 6.3.3 for production data verification and audit trail requirements.

For delivery point measurement, hydrocarbon liquid volume must be determined to 2 decimal places and rounded to 1 decimal place for monthly reporting. If there is more than one volume determination within the month at a reporting point, the volumes determined to 2 decimal places must be totalled prior to the total being rounded to 1 decimal place for reporting purposes.

10.1.2 Temperature Correction Requirements

All delivery point measurement of hydrocarbons and emulsions requires temperature correction to 15°C (see Section 6.3.1.1). See Section 6.3.2.1 for temperature determination requirements. Composite meter factors are not acceptable for delivery point measurements.

The temperature correction factor (CTL) must be determined in accordance with the API MPMS, Chapter 11.1. LPG must follow the applicable API Table 54E or an equivalent industry standard.

10.1.3 Pressure Correction Requirements

The pressure correction factor (CPL) must be determined in accordance with API MPMS, Chapter 11, and is required only for LACT applications.

10.2 General Trucked Liquid Measurement, Accounting, and Reporting Requirements for Various Facility Types

10.2.1 Conventional Oil Batteries

For trucked oil/emulsion production into an oil battery, delivery point measurement is required for the total liquid volume. If there is a mixture of trucked-in production and prorated production within the same battery, the exception criteria in Section 5.5.1.2 must be met or ERCB site-specific approval must be obtained.

For condensate trucked into an oil battery, delivery point measurement is required for the total liquid volume. The requirements in Section 6.6 must be met.

For any oil battery, the trucked-out liquid is measured at the delivery point (located at the receiving facility), and the oil volume determined at the receiving facility must be used as the delivering battery's oil disposition. Proper measurement must be set up at the receipt point only, except for load oil delivery from a facility to well(s). In this case, delivery point measurement is required at the loading facility. If there are emergencies or problems at the receipt point, the origin measurement may be used, but only as a temporary solution.

If clean oil from a battery is delivered into an oil pipeline via a LACT meter and that same LACT facility also receives clean trucked oil from other sources, a terminal licence and PRA code must be obtained so that the Alberta Petroleum Marketing Commission (APMC) can allocate royalty and oil equalizations to all the oil receipt volumes.

10.2.2 Custom Treating Facilities

The measurement requirements are the same as for trucking into a conventional oil battery above, but the accounting and reporting must follow the requirements in *Directive 048: Monthly Custom Treating Plant Statement*.

10.2.3 Pipeline Terminals

At the pipeline terminals that receive either pipelined and/or trucked clean oil, the receipt meter or weigh scale measurement is considered as LACT measurement. That is, there is no proration/allocation from the pipeline disposition volumes to the receipt volumes. However, the downstream pipeline operator may allocate to the shippers the imbalance (generally less than 1%) on their pipeline system according to their contractual requirements.

10.2.4   Clean Oil Terminals

Clean oil terminals are those that receive trucked clean oil only; the receipt meter is considered as a delivery point. That is, there will be proration/allocation from the terminal LACT disposition volumes to the receipt volumes for that month.

Exception

Volumetric allocation of the monthly LACT volumes to the monthly truck receipt volume is not required at a clean oil terminal without ERCB site-specific approvals if all of the following criteria are met:

- The meter factor for each delivery point meter or the weigh scale accuracy verification must not deviate from the prior factor or verification by more than 0.5%.

- Any deviation over 0.5% must be investigated and rectified, and allocation for the previous month(s) disposition volumes to the receipt volumes is required.

- The monthly measurement difference between the receipts and dispositions after including allowances for blending shrinkage, if applicable, and the inventory changes at the clean oil terminal must be less than or equal to 0.5%.

The licensee must revert to allocating monthly pipeline LACT volumes to the receipt volumes if any of the above criteria are not met.

10.2.5   Gas Plants, Gas Batteries, and Gas Gathering Systems

For gas systems receiving trucked liquid, the measurement requirements are the same as for trucking liquid into a conventional oil battery.

10.2.6   Water Injection/Disposal Facilities

For water trucked into an injection or disposal facility, delivery point measurement accuracy is not required. See Section 1.7 for facility accuracy requirements.

10.2.7   Waste Processing Facilities

A waste processing facility handles volumes of waste generated in the upstream petroleum industry. However many ERCB-approved waste facilities have an integrated custom treating facility designated for processing oil/water emulsions extracted from the solids during waste processing. In addition, oil/water emulsions from other batteries are trucked in and measured independently from the waste oil/water emulsions, and both streams are processed through the same treating facilities. The total waste stream disposition to the custom treater (CT) must have emulsion volume and S&W determinations in order to properly allocate the clean oil and water volumes back to the other received emulsions. Therefore, delivery point measurement is required at the receipt point of non-waste truck unloading and at the total waste oil/emulsion delivery point to the CT for further processing, such as in a treater, where it is commingled with other oil/emulsion from other sources.

There are also injection/disposal facilities that receive other liquids, such as waste streams going into subsurface caverns for disposal. Waste liquids for disposal require measurement accuracy similar to disposal of produced water.

See *Directive 058: Oilfield Waste Management Requirements for the Upstream Petroleum Industry* for the requirements for waste stream measurement, accounting, and reporting.

10.2.8   Heavy Oil and Crude Bitumen Facilities

To meet heavy oil and crude bitumen trucked production delivery point measurement requirements, the licensee must use an appropriate method based on the fluid characteristics, such as viscosity, temperature, and sand content of the load. Generally, delivery point measurement is performed by using weigh scales or tank gauging with sampling to determine the S&W and/or density. Meters are used only when there are minimal or no solids present in the oil/emulsion, similar to trucking into a conventional oil battery.

10.3   Design and Installation of Measurement Systems

Delivery point measurement is required for most trucked fluids delivery/receipt except as mentioned above. The gross volume must be measured through a system consisting of inlet tank gauging, inlet meter, or weigh scale. Gauge boards must not be used for delivery point measurement.

Truck ticket estimates (i.e., volume estimates determined using the truck tank load indicator completed by the trucker or trucking company for bill of lading/transportation of dangerous goods purposes) are not considered as measurement for the purpose of well or facility volume measurement. Therefore, truck ticket estimates must not be used for determining volumes unless the requirements in Section 10.3.4 are met.

10.3.1   Tankage

Inlet tankage used for the purpose of truck volume measurement must be of sufficiently small diameter in relation to load volume to facilitate gauging accuracy to within the Standard of Accuracy requirements in Section 1. The sizing of the maximum diameter of a tank for measurement purpose and other tank gauging requirements are in Section 6.3.1.2a. An exemption from this requirement is granted only during seasonal road ban periods when reduced truck loads are mandated by weight restrictions.

10.3.2   Meters

Meter selection and sizing for truck unloading must be appropriate for the intended range of service, having regard for the range of flow rates and liquid characteristics (e.g., density, viscosity, temperature) to be encountered. To prevent exceeding the range of a meter, in some cases it may be necessary to use a valve to control the truck unloading rate. See Figures 10.1 and 10.2 for examples of system setup.

Turbine meters are typically not suitable for viscous fluids and therefore are not recommended for unloading crude oil.

When metering devices for the purpose of measuring truck delivery/receipt volumes are installed, the following must also be installed:

- sample point
- air eliminator
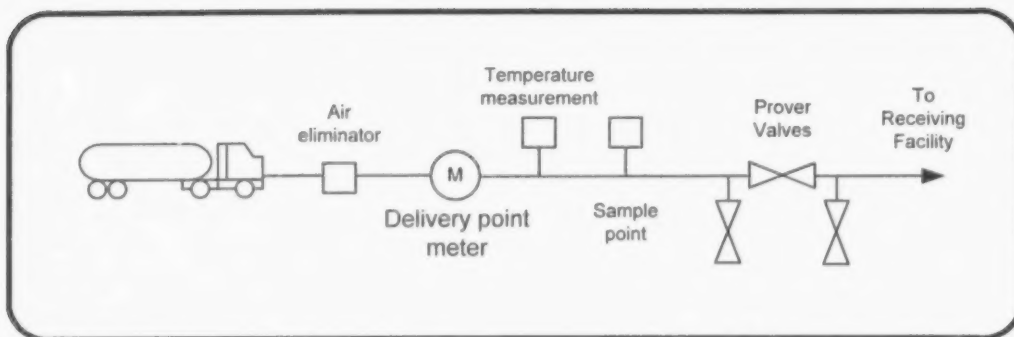
Figure 10.1. Typical truck unloading measurement

For some types of meters and applications, a strainer and a back pressure control system are required. See Figure 10.2.
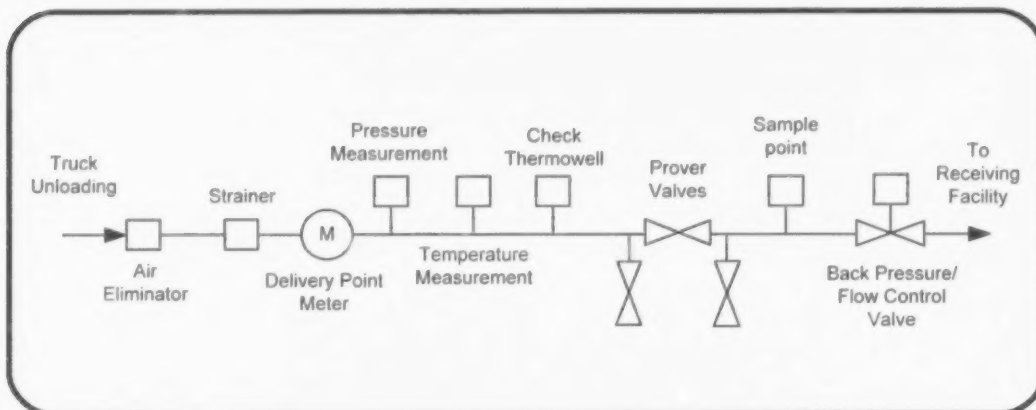
Figure 10.2. Typical truck unloading measurement for high vapour pressure liquids

Additional requirements for clean oil and pipeline terminals:

1) For mechanical automatic temperature compensators without gravity selection (ATC) or with gravity selection (ATG):

   • For new applications, mechanical ATC and ATG must not be used. All existing ATC and ATG are grandfathered at their existing applications and must not be relocated or reused for other applications.

   • The difference between actual density and compensation density must be less than 40 kg/m^3.

   • Product temperature must be between –10°C and +40°C excluding LPGs.

   • The compensation density (mechanically set density or user-entered density for electronic flow computers) must be a volume weighted average of the expected receipt volumes. When product temperatures exceed 40°C, it may be necessary to reduce the allowable density difference to maintain a 0.5% uncertainty.

- Temperature compensation devices must be designed for the actual range of operating temperatures observed. If product temperatures exceed 40°C, it may be necessary to reduce the allowable density difference to maintain a 0.5% uncertainty.

2) For positive displacement meters, temperature compensation can be achieved by using mechanical ATC or ATG (see item 1 above) or through electronics using flow computers or by manual temperature compensation. When manual temperature compensation is used, an average temperature based on an initial flowing temperature (taken 2 to 5 minutes after the start of unloading) and a final flowing temperature (taken 2 to 5 minutes prior to the end of unloading) must be used.

3) A thermowell located either within the meter body or not more than 20 pipe diameters downstream of the meter without restrictions in between is required for temperature determination.

4) For applications where there is a potential for the product to flash, a pressure indicator downstream of the meter to check backpressure is required.

10.3.3 Weigh Scales

Weigh scales for the purpose of delivery point measurement must be verified in accordance with Section 2.14. For sampling points and methods, see Section 6.4.2. Systems employing weigh scales must also provide for determination of density of oil and water in accordance with one of the following:

1) API MPMS, Chapter 9: Density Determination Using Hydrometer,

2) precision laboratory method – ASTM method, or

3) on-line densitometer (Coriolis, nuclear device, etc.).

To maintain an uncertainty of 0.5% or less, the net weight of the payload must not be less than 40% of the gross vehicle weight and the net weight must not be less than 6500 kg. An exemption from this requirement is granted only during seasonal road ban periods when reduced truck loads are mandated by weight restrictions.

10.3.4 Exceptions

10.3.4.1 Truck-Mounted Level Gauges and Truck-Mounted Meters

Truck gauge level indicators and truck-mounted meters are considered to have met the requirement for low-accuracy measurement with an overall uncertainty of ±1% or less if the following criteria are met. These units can be used for trucked-in delivery point measurement to proration oil batteries or group oil batteries if all of the following requirements are met:

- the battery receives not more than 100 m^3 of trucked-in oil per day;

- the maximum percentage of trucked-in oil to any battery is 10% of the monthly total battery oil production volume;

- the gauges or meters are verified annually and if not within ±1% accuracy they are repaired and reproved;

- the product temperature is determined to within 1°C (see Section 10.3.2, item 2); and

- the truck gauge levels or meters are initially set by calibrating to a master meter or provers with a demonstrated uncertainty of not more than 0.2%.

Additional criteria for truck-mounted level gauges:

- the stated depth of liquid is within 12.7 mm of a known gauge level marker if used;

- the depth of liquid is determined while the tank trailer is level to within 150 mm over its length; and

- the minimum load on the trailer is more than 65% of full load.

10.3.4.2 Truck Tickets and Lease Tank Gauging

Truck ticket volumes uncorrected for temperature are not acceptable for delivery point measurement of trucked liquid except if the fluid transfers are between unitized or common equity facilities with common royalty; then the temperature correction estimates may be used. The truck ticket must be based on a low-accuracy measurement requirement with an overall uncertainty of ±1% or less of trucked liquid, such as lease tank gauging at the battery sending the liquid production or truck-mounted meter, for determining inlet volumes at a proration battery if certain situations exist. The S&W% and corrected opening and closing readings must be on the ticket or available on a summary sheet for ERCB audit purposes. An individual truck load should be recorded on its own ticket.

The ERCB may accept low-accuracy measurement with an overall uncertainty of ±1% or less for trucked liquid production at a proration battery if

- trucked production is temporary, pending battery consolidation within one year or less;

- individual well oil volumes being trucked are less than 2.0 m^3/day (see Section 10.3.5);

- the crude oil volume receipt (net of water) is 5% or less of the total receiving battery oil production; and

- truck-mounted meters used for low-accuracy measurement with an overall uncertainty of ±1% or less are proved in accordance with the requirements in Section 2.

10.3.5 Load Fluids

Load fluids at a minimum must be measured using devices that meet the requirement for low-accuracy measurement with an overall uncertainty of ±1% either at the source (loading) location or at the delivery point.

Reporting of load fluid on the PRA is limited to oil-based and/or water-based fluid(s) injected during preproduction well stimulation or postproduction activities. Only the load fluid product codes OIL, COND, or WATER can be reported. Well drilling fluids must not be reported on the PRA as load fluids.

Load fluids are to be reported at the well level except when in an SE Alberta shallow gas battery, since there is no requirement to measure and report water production at this type of well. The load fluid reporting then can be done at the battery level.

See the *Directive 007: Volumetric and Infrastructure Requirements* (December 2007) on-line supplement Appendix 8 for more reporting procedures.

10.3.6 Split Loads

A split load is defined as existing when a truck takes on partial loads from more than one well or battery in a single trip or when load oil is delivered to more than one receipt point or wells.

Allowed	- Single-well oil battery delivering to other facilities
	- Gas wells with condensate-water tanks and production less than 2.0 m^3 of total liquids per day
	- Blending of heavy oil and condensate
	- Load oil for well servicing only; load up from a single source only
Not allowed	- Multiwell batteries delivering to other facilities other than load oil
	- Gas wells with production greater than 2.0 m^3 of total liquids per day
Requirements	Densities must be "similar" (within 40 kg/m^3). If they are not, blending tables are required to calculate shrinkage. The shrinkage volume is to be prorated back to each battery on a volumetric basis.
Measurement	Volume from each well or facility must be measured at the time of loading onto the truck (or off loading from the truck for load oil) by one of the methods below:
	1) gauging the battery lease tank;
	2) gauging the truck tank (not allowed for density difference over 40 kg/m^3 for any oils or emulsions); or
	3) truck-mounted meter/gauge that meets low-accuracy measurement and is proved at least annually.
	Calibrated gauge tables are required for methods (1) and (2) above.
Sampling	Fluid from each single-well oil battery must be sampled to determine the S&W and the oil/water volumes. The truck driver is to collect the samples by taking at least three well-spaced grab samples during the loading period (see Section 6.4.2).
	For load oil, the S&W must be determined at the loading source.
Records	The truck tickets must show the individual load volumes, as well as the total volume at delivery (receipt) point, supported by opening and closing gauge or meter readings.
Accounting	For battery emulsions, the total load is to be measured and sampled at the receiving location and prorated to each of the wells based on the measured loading volumes and S&W from each of the wells.
	For load oil, the initial volume must be measured at the loading source and prorated to each delivery point based on the measured volume delivered to each well.

10.4 Sampling and Analysis

For trucked-in hydrocarbons and emulsions receipt/delivery, a truck thief sampler or a proportional sampler may be used to obtain a sample from the truck tank (see Section 6.4.2). In some cases spot (grab) samples may be used to obtain the sample from the off-load/load line. Automatic sampling methods are preferred; however, manual or tank sampling systems may also be allowed, as discussed below.

The frequency of sampling or readings must be sufficient to ensure that a representative sample of the entire truck volume is obtained. Consideration must be given to both conditioning the flow stream and locating a probe or sampler. Flow conditioning to ensure turbulent mixing can be achieved through velocity control, piping configurations, or introduction of a mixing element upstream of the sample point. A sample probe is required for truck delivery point sampling unless there is an in-line product analyzer or the sampling is incorporated as part of the measurement system. A mid-pipe probe location should be used for accurate sampling. (Also see Section 8.2.1.)

The licensee must choose the sampling methodology based on emulsion characteristics, stratification, and S&W consistency of each load to obtain a representative sample. API MPMS, Chapter 8.1, Section 8, provides further information on manual sampling procedures.

10.4.1 Automatic Sampling

Automatic sampling is typically conducted through the use of proportional samplers. If automatic sampling procedures are used, a manual procedure should also be in place for use when the automatic system is out of service or for intermittent verification of the automatic system reading. For more information, API MPMS, Chapter 8.2, Sections 7 to 15, provide further details on flow conditioning, probe location, and sampling frequency.

Other requirements for automatic sampling:

- Containers made of suitable material for handling and storage of the sample must be used. Container lids must be vapour tight.

- All sample containers must be cleaned and dried prior to collection of the next sample.

- Sample containers must allow adequate room for expansion and content mixing, taking into consideration the temperature of the liquid at the time of filling.

- The sample containers must be housed in a secured enclosure to prevent any tampering with the sample.

- Sample lines must be as short as practical and sloped downward to reduce the possibility of plugging up the sample line.

10.4.2 Manual Spot (Grab) Sampling

Manual spot (grab) sampling may be acceptable in situations involving a tight emulsion with less than 0.5% S&W in the truck by taking three well-spaced grab samples during the unloading period (see Section 6.4.2). Grab samples are not acceptable when there is stratification of S&W within the truck.

The use of manual sampling techniques, such as truck tank thieving (full height or intermittent), may also be acceptable. However, in the presence of stratification, one unit of height at the bottom of the truck tank represents a significantly lesser volume than the same unit of height at the midpoint of the truck tank because of the shape of the tank. The resulting S&W from a full-height core thief therefore may not be representative of the entire load. Then multiple grab samples is to be used.

Lease tank thieving is subject to similar stratification limitations (excluding the non-uniformity of the tank). These concerns can be reduced by locating any water-emulsion interface and obtaining bottom, middle, and top samples of the emulsion to determine the average water cut of the emulsion. However, lease tank thieving requires dedicated tankage for each load received or delivered to avoid mixing of product between deliveries.

Visual estimates or estimates based on changing off-load pump speeds must not be used for free water volume determination.

10.4.3 S&W Determination

The licensee must select the most appropriate method for determining the S&W. See Section 6.4.3 and Appendix 4.

10.4.4 Density Determination

Truck load sample density determination at 15°C must be conducted at least annually or more frequently if there are changes in the reservoir conditions. Density of the load may be determined by one of the following methods:

1) Truck load samples may be collected from the receiving point and sent to an independent laboratory for analysis to determine density of the hydrocarbon phase and the water phase (if required). The density found in this analysis must be applied to all hydrocarbon liquids coming from the specific facility.

2) Truck load samples or samples from automatic samplers may be tested for density as outlined in API-MPMS, Chapter 9.1: Hydrometer Test Method for Density, Relative Density (Specific Density).

There are two methods of density determination using densitometers or other density analyzers: dynamic and static. The user must ensure that the selected method is suitable for the application and determines the density of the entire sample, which could include emulsion and free water.

- The dynamic method can be conducted using an on-line densitometer or a Coriolis meter.

- Static determination can be conducted by either the hydrometer method (see API MPMS, Chapter 9) or the precision densitometer method (see ASTM D4052).

In applications where the truck volumes have an S&W greater than 1%, density determination at 15°C of an emulsion sample is difficult, as there are two different thermal corrections to be applied, one for the water and one for the oil.

There are two options available:

- The first is to determine the sample density using a precision densitometer that has its measuring cell at 15°C. No further corrections are required.

- The second is to separately predetermine the density at 15°C of the water and the oil. When using this option, the emulsion density is calculated by applying the S&W cut to the density of each component. The calculation is

$$\rho_{emulsion} = \left(\rho_{oil} \times \left(100 - \%S \& W\right)\right) + \left(\rho_{water} \times \%S \& W\right)$$

where

$\rho_{emulsion}$ is the calculated density of the emulsion at 15°C

ρ_{oil} is the density of the oil portion at 15°C

ρ_{water} is the density of the water portion at 15°C

10.5 Volume Determination

10.5.1 Tank Gauging

Tank gauging procedures are detailed in Section 6.3.1.2. The starting and closing levels measured are then converted to volume through the use of gauge tables (supplied by the tank manufacturer), which have been calculated using measurements of the tank. The difference between the closing and opening volumes is the measured volume. If the tank is used for delivery point measurement, the temperature and density of the tank contents must be taken in order to correct the indicated volume to standard conditions before determining the volume difference.

10.5.2 Weigh Scales

The procedure for determining the volume of liquid on a truck using a scale is to weigh the truck before and after loading or unloading and determine the difference to obtain the net weight. The entire load must be weighed at a time—i.e., split weighing, whereby the truck is weighed after unloading a portion of its load to obtain the weight of the unloaded portion is not permitted.

To determine the density of the load, an on-line densitometer maybe used or a representative sample must be obtained and the density and temperature measured with a hydrometer and thermometer respectively. The observed density must be corrected to 15°C.

The net weight determined during the weighing process divided by the sample density at 15°C results in the net volume of the load prior to deductions for S&W.

10.5.3 Meters

Metered volumes must be determined in accordance with Section 6.3.2.

11 Acid Gas and Sulphur Measurement

This section presents the base requirements and exceptions for acid gas and sulphur measurements at processing plants and injection facilities in the upstream oil and gas industry that are used in determining volumes for reporting to the ERCB. *S-30 Monthly Gas Processing Plant Sulphur Balance Report* requirements are also included, with instructions provided in Section 11.6.

In a gas processing plant where sour gas is processed, most of the acidic portion of the gas must be removed from the gas stream (sweetening) in order to produce a saleable pipeline-quality gas product. However, in the process of removing the acidic portion of the sour gas, acid gas, which consists mainly of H_2S and CO_2, is generated and must be disposed of in an environmentally and economically acceptable way, such as by elemental sulphur production, acid gas injection, or acid gas flaring.

11.1 General Requirements

The sour gas plant inlet and acid gas streams must be measured and reported. The inlet sour gas stream volume including GEV of condensate, sour gas in solution in water, the sulphur disposition tonnage, and the sulphur balance must be reported on a monthly basis on the S-30 report if the plant is approved with a sulphur inlet of more than 1 tonne per day (t/d) to the ERCB. See Table 11.1 for the monthly S-30 sulphur balance requirement. For sour gas plants with less than 1 t/d of approved sulphur inlet, the S-30 report must be submitted to Alberta Environment at its required timing, with the exception of grandfathered plants that would still be required to submit S-30 reports to the ERCB until December 31, 2016. For plants that are licensed as "sweet" but use a sweetening process to strip out excess CO_2, the reporting of the acid gas (CO_2) volume must be the same as sour plants under 1 t/d.

Table 11.1. Monthly S-30 Sulphur Balance Requirement

Monthly average actual sulphur inlet (t/d)	Sulphur balance %
< 1	20%
≥ 1	5%

In accordance with *Interim Directive (ID) 2001-03: Sulphur Recovery Guidelines for the Province of Alberta*, other upstream oil and gas facilities with sulphur emissions greater than 1 t/d that are not required to submit S-30 reports must maintain daily sulphur balance records and calendar quarter-year recovery calculations. These records must be available for inspection or audit at the request of the ERCB.

The acid gas from the sweetening process is generally saturated with water vapour (wet). This water vapour portion must be subtracted from the saturated acid gas to obtain the dry volume without water vapour. (See Section 11.2.)

11.2 Acid Gas Measurement

The quantity of acid gas going to sulphur plants, to compression and injection, or to flaring is generally measured at a low pressure of 50 to 110 kPag; therefore, the orifice meter, or any other meter, must be appropriately sized and well maintained to achieve accurate measurement.

Acid gas is saturated with water vapour, which represents a significant portion of the total gas measured. The amount of water vapour varies significantly with the temperature of the reflux drum. Therefore, it must have continuous temperature measurement to calculate the correct

acid gas volume (see below). The gas density must also include the water content, and the meter coefficient must include a factor to exclude the water vapour content of the gas in the final volume computation for reporting purposes. The accuracy of the gas relative density factor and water content determination must be periodically verified to ensure that acid gas measurement uncertainty is within tolerance.

11.2.1 Determining Acid Gas on a Dry Basis

For ideal gases, the total vapour pressure of a system containing several components is the sum of the vapour pressure of the individual components at the temperature of the system.

The component's vapour pressure percentage of the total pressure of a system is equal to the volume percentage of that component in the system. The reflux drum is the vessel in which the acid gas separates from the sweetening solution. The amount of water vapour in the acid gas leaving the reflux drum is a function of the temperature and the absolute pressure in the reflux drum.

Calculating Acid Gas Flow Rate

The recommended calculation method for the acid gas flow rate is as follows:[1]

Step 1: Determine the percentage of water vapour in the acid gas on the basis of the ratio of vapour pressure of water to total pressure in the reflux drum at the reflux drum temperature.

Step 2: Convert the acid gas composition from dry basis to wet basis at reflux drum pressure and temperature, and determine the acid gas relative density and compressibility factor on a wet basis at meter pressure and temperature.

Step 3: Calculate the acid gas and water vapour flow rate corrected from actual flowing pressure and temperature to standard conditions of 101.325 kPa(a) and 15°C.

Step 4: The volume calculated in step 3 contains water vapour in the percentage determined in step 1 and must be converted to dry basis volume for reporting purposes. A **correction factor (CF)** has to be applied to correct the acid gas flow from a wet to a dry basis.

CF = (100.00 – % H_2O in acid gas)/100

Dry acid gas flow rate = CF x flow rate calculated in Step 3

The H_2S content of the acid gas is the dry basis acid gas flow times the % H_2S/100 in the acid gas on a dry basis.

Calculating Vapour Pressure of Water

The formula for determining the vapour pressure of water[2] is

$\log P = A - B/(C + T)$

where   P = water vapour pressure in mm of mercury
         A = 8.10765
         B = 1750.280
         C = 235

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[1] Wichert, E., "Water content affects low pressure, acid-gas metering," *Oil & Gas Journal*, January 2, 2006, pp. 44-46.

[2] The vapour pressure of water at a certain temperature can also be obtained from the GPSA *Engineering Data Book*, SI Units version, 12th edition, 2004 or subsequent versions, Figures 24-36.

$$T = \text{temperature of acid gas in reflux drum, °C}$$

The direct formula for determining the vapour pressure of water in kPa(a) is thus

$$P_{H2O} = 0.13332 * 10^{(8.10765 - 1750.280/(235 + T))}$$

where P_{H2O} = water vapour pressure in kPa(a) at T°C

% H_2O in the acid gas = 100 % * $P_{H2O}/(P_{RD} + P_{atm})$

where P_{RD} = reflux drum pressure, kPag

P_{atm} = atmospheric pressure, kPa(a)

This method for estimating the water vapour content of acid gas replaces the method contained in Section B-2 of *Directive 046: Production Audit Handbook*. There is no grandfathering for using the methodology in *Directive 046*. Noncompliance will not be assessed until after May 1, 2009, to allow industry sufficient time to implement the changeover of methodology.

Converting Acid Gas Calculation from Dry to Wet Basis

An example acid gas conversion calculation from dry to wet basis with the meter installed upstream of the back-pressure regulator of the reflux drum is provided below:

A. Reflux drum data

Reflux drum temperature = 40°C
Reflux drum pressure = 70 kPag
Atmospheric pressure = 95.0 kPa(a)

If the meter is installed upstream of the back-pressure regulator of the reflux drum, the upstream pressure and the temperature of the meter run may be used as the reflux drum pressure and temperature.

B. Acid gas components on a dry basis from acid gas analysis:

H_2S = 65%; CO_2 = 33.5%; C_1 = 1.2%; C_2 = 0.3%

C. Calculate the percentage of components, including water vapour, on a wet basis:

Percentage of water vapour = (100 % * Vapour pressure of water at 40°C) / (Reflux drum gauge pressure + atmospheric pressure)

$$P_{H2O} = 0.13332 \times 10^{(8.10765 - 1750.280/(235 + 40))}$$
$$= 7.377 \text{ kPa(a)}$$

(Vapour pressure of water at 40°C is 7.384 kPa(a), from the Saturated Steam Table in the Thermodynamics section of the GPSA *SI Engineering Data Book*, Figures 24-36.)

Percentage of water vapour = 7.377/(70 + 95) * 100% = 4.47%

Enter into column 2 (see the following table) and normalize.

Calculation of relative density (RD) on wet basis

Comp.	Col. 1 Dry basis (%)	Col. 2 Wet basis (%)	Col. 3 Molar mass (kg/kmol)[1]	Col. 4 (Col. 1 * Col. 3) /100	Col. 5 (Col. 2 * Col. 3) /100
H_2S	65.00	62.09	34.082	22.153	21.162
CO_2	33.50	32.00	44.010	14.743	14.083
C_1	1.20	1.15	16.042	0.193	0.184
C_2	0.30	0.29	30.069	0.090	0.087
H_2O	0.00	4.47	18.0153	0.000	0.805
Total	100.00	100.00		37.179	36.321

[1] Molar mass of air = 28.9586 kg/kmol (GPSA *Engineering Data Book*, 2004 or later editions, Figure 23-2, or GPA-2145).

From column 4, ideal gas RD, dry basis = 37.179/28.9586 = 1.284.

From column 5, RD wet basis = 36.321/28.9586 = 1.254 (this RD is to be used in the flow calculation for acid gas volumes).

11.2.1.1 Difference Between the Acid Gas Volume on a Wet Basis and on a Dry Basis

An example calculation is presented to show the difference in the results of the acid gas flow rate and the sulphur content of acid gas using dry versus wet basis metering. The example data for the meter run and assumed conditions are as follows:

- Orifice meter diameter: 154.051 mm
- Orifice plate diameter: 76.200 mm
- Meter upstream pressure: 70.00 kPag
- Differential pressure: 10.00 kPa
- Meter temperature: 40°C
- Atmospheric pressure: 95.0 kPa
- Acid gas composition: as per above table

Results with AGA # 3 1992 or later method:

- Flow rate, dry basis without accounting for moisture content = 33.126 $10^3 \, m^3$/d
- Sulphur content = 33.126 x 65/100 * 1.35592 = 29.20 t/d
- Flow rate, wet basis = 33.499 $10^3 \, m^3$/d, containing 4.47 vol % H_2O
- Flow rate, wet basis converted to dry basis = 33.499 * (100 – 4.47)/100
= 32.002 $10^3 \, m^3$/d dry acid gas equivalent

This volume, 32.0 $10^3 \, m^3$, is to be reported as "Acid Gas" on the monthly volumetric submission.

An example for percentage difference in acid gas volume between dry and wet basis:

- Percentage difference in flow rate = (33.126 – 32.002) * 100 %/32.002 = 3.51%
- Sulphur content = 32.002 * 65/100 * 1.35592 = 28.21 t/d
- Difference in calculated sulphur balance between dry and wet basis metering
= 29.20 – 28.21 = 0.99 t/d
- Percentage difference = 0.99 x 100/28.21 = 3.51%

Thus, if the moisture content in the metering of the acid gas in this example were ignored (dry basis taken as wet basis), the reported acid gas flow and sulphur content in the acid gas leaving the reflux drum would be 3.51% higher than the correct value.

This method of estimating the water vapour content is valid when the gas is in contact with water in a low-pressure vessel, such as in the reflux drum. The method does not apply to low-pressure gas, such as in a flare line, when the flared gas originates from a high-pressure vessel.

The table below summarizes the above figures and also provides the results that are obtained by the 1985 AGA # 3 Report method, using Wichert-Aziz (W-A) compressibility factors.

Item	AGA #3, post-1992		AGA #3, 1985, W-A Z factors*	
	Dry basis	Wet basis	Dry basis	Wet basis
Z factor at St'd P and T	0.992 848	0.991 999	0.993 037	0.992 788
Z factor at Meter P,T	0.991 002	0.990 030	0.991 007	0.990 674
Flow rate, 10³ m³/d	33.126	33.499	33.096	33.481
Corrected to dry gas	-	32.002	-	31.984
% difference	-	3.51	-	3.48
Sulphur flow, t/d	29.20	28.21	29.17	28.19
% difference, t/d		3.51		3.48

*Z factors by Wichert-Aziz method, including water content in wet gas.

11.2.1.2 Calculation Method of Water Content if Meter Located Downstream of Back-Pressure Valve of Reflux Drum

The water content in the acid gas is a function of the pressure and temperature of the reflux drum. If the acid gas meter is located downstream of the back-pressure regulator of the reflux drum, both the pressure and the temperature of the meter will be somewhat lower than the pressure and temperature of the reflux drum. Under these conditions, it is still necessary to determine the water vapour content of the acid gas stream at the reflux drum pressure and temperature, as shown above, to correctly calculate the acid gas flow rate.

The reflux drum pressure must be recorded for the correct calculation of the water vapour content of the acid gas. The reflux drum temperature should also be used to estimate the water content. However, since the flow data from the meter include the temperature at the meter run, the reflux drum temperature can be estimated on the basis of the meter temperature, as follows:

$$T = T_m + 2.28 - 2.28 * P_2/(P_{RD} + P_{atm})$$

where P_2 = the downstream meter tap pressure, kPa(a)

T_m = temperature downstream of the orifice plate

Having estimated the temperature at the reflux drum from the temperature downstream of the orifice plate, the vapour pressure of the water can be calculated as above. The percentage of water vapour in the acid gas can then be determined using the reflux drum pressure, and the same procedure as outlined in the above example can be used to calculate the acid gas flow rate.

11.2.1.3 Effect of Compression and Cooling of Acid Gas

In the situation of acid gas compression and injection, the acid gas flow rate may in some instances be metered after one or more stages of compression and cooling. This will remove a sufficient amount of water so that the remaining water vapour in the compressed and cooled

acid gas will have little effect on the acid gas metering. In such a situation, it is not necessary to include the effect of water vapour in the metering of the acid gas.

11.3 Sulphur Measurement and Pit Volume Determination

11.3.1 Sulphur Pit Volume/Tonnage Determination

When pit gauging is used to determine a liquid sulphur volume, the gauging procedures must be conducted in accordance with the following:

- The operator must ensure that the gauge/strapping table used to convert the gauge level to a liquid volume is specific for the pit being gauged.

- Pit gauging should be used for inventory determination only and must not be used for delivery point measurement.

- All dip sticks and electronic level devices must have a minimum resolution of 6 mm.

- It is acceptable to have one reading per determination.

- The sulphur density at pit temperature is obtained from Figure 11.1.[3]

The general formula for determining the produced sulphur tonnage is as follows:

Sulphur tonnage = Gauge reading x CF x Sulphur density

where CF = Pit gauge/strapping table conversion factor

11.3.2 Sulphur Measurement

For sulphur sales/delivery point measurement using meters, see Section 6.3.1.1. These meters must be kept at a temperature so that the molten sulphur will not solidify when there is no flow.

For sulphur sales/delivery point measurement using a weigh scale, see Section 2.13.

For daily sulphur production measurement using pit level gauging, two pits are required, one for production and the other for withdrawal using level measurement. The daily sulphur production tonnage should be adjusted by the total monthly disposition at the end of the month (see Section 11.3.2.1).

11.3.2.1 Exceptions

For daily sulphur production volume determination, if there is only one pit in place in an existing plant and sulphur is being withdrawn without measurement, the operator may use the measured acid gas volume on a dry basis, provided that there is a continuous acid gas sampling device, such as a gas chromatograph, to calculate the sulphur tonnage entering the sulphur plant. The daily sulphur production can then be calculated using the following formula:

Estimated daily sulphur production (t) = Daily acid gas inlet (t) – Daily incineration (t) –
Daily flared (t) – Others if applicable (t)

The estimated daily sulphur production tonnage must be adjusted by the total monthly disposition at the end of the month by calculating a proration factor and applying that to all estimated daily production tonnage:

---

[3] Tullen, W. N., *The Sulphur Data Book*, New York: McGraw-Hill Book Company, Inc., 1954, p.17.

Sulphur proration factor (S_{pf}) = Total monthly sulphur disposition tonnage (including inventory changes) + Total estimated daily sulphur production tonnage

Actual daily sulphur production (t) = Estimated daily sulphur production (t) * S_{pf}

The "actual daily sulphur production" is the daily production tonnage to be reported on the S-30 Monthly Gas Processing Plant Sulphur Balance Report.
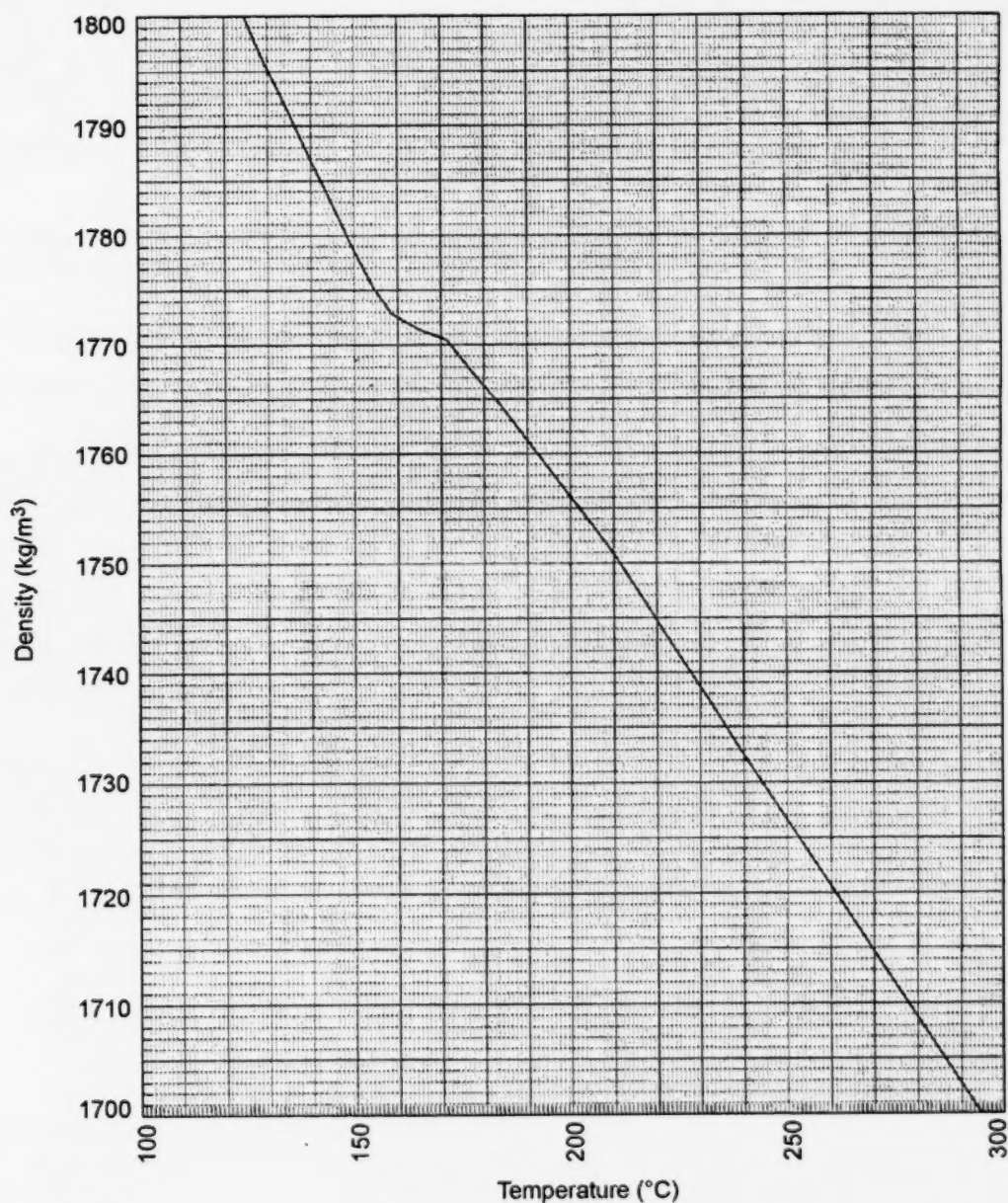
Figure 11.1. Liquid sulphur density vs. temperature

11.4 Sulphur Balance Calculation for Sour Gas Processing Plants

When sour gas is produced to a sour gas treating plant, it always enters the plant through a plant inlet separator. A liquid water phase is usually present with the sour gas, and in many instances a liquid hydrocarbon phase can also be produced into the separator with the gas and water. In such situations, all three phases will contain some H_2S in different proportions.

All of the H_2S entering the plant in the different fluids will also exit the plant by one means or another. The balance is an important part of checking to ensure that all streams are accounted for and reported. The S-30 reports a monthly balance between the accounting of the mass of sulphur entering and leaving the plant for sour gas plants with an approved sulphur inlet rate greater than 1 t/d. To achieve this objective, certain minimum measurements of concentrations and flow volumes must be made by the plant operator. This section provides the minimum requirements to achieve the desired goal.

If a monthly balance in Table 11.1 in Section 11.1 cannot be achieved on a regular basis, the operator must implement appropriate measures to ensure that the required plant-wide balance is achieved. Appropriate measures include, but are not limited to,

- improving the acid gas, inlet gas, flare, and sulphur measurement systems;
- installing a continuous gas analyzer on the gas stream of the plant inlet separator and/or on the outlet gas stream of the reflux drum for acid gas flaring plants;
- installing a proportional sampler on the gas stream of the plant inlet separator; and
- improving the methodology for determining sulphur content in inlet condensate and water.

11.4.1 Overview of Plant Inlet and Outlet Points for H_2S

Figure 11.2 illustrates the paths by which H_2S enters the sour gas plant and by which method it can exit from the plant.

The H_2S entering the plant in the gas, condensate, and water has to be accounted for on the S-30 report in terms of tonnes of elemental sulphur. Since the flow volumes of the three types of fluid streams out of the inlet separator are required to be measured, it becomes a simple task to account for the amount of H_2S entering the plant by determining the H_2S concentration in each stream.

11.4.2 Determining H_2S Contents

11.4.2.1 Determining H_2S in Sour Gas

See Section 8.2 for the determination of H_2S concentration in the inlet gas stream.

11.4.2.2 Determining the Concentration of H_2S in Condensate

Condensate associated with sour gas will contain some H_2S. The physical relationship between the concentration of H_2S in the gas and in the condensate depends on the composition of the gas and the condensate and the pressure and temperature in the plant inlet separator. The concentration of the H_2S in the condensate is usually determined in a laboratory on condensate samples obtained from the inlet separator.
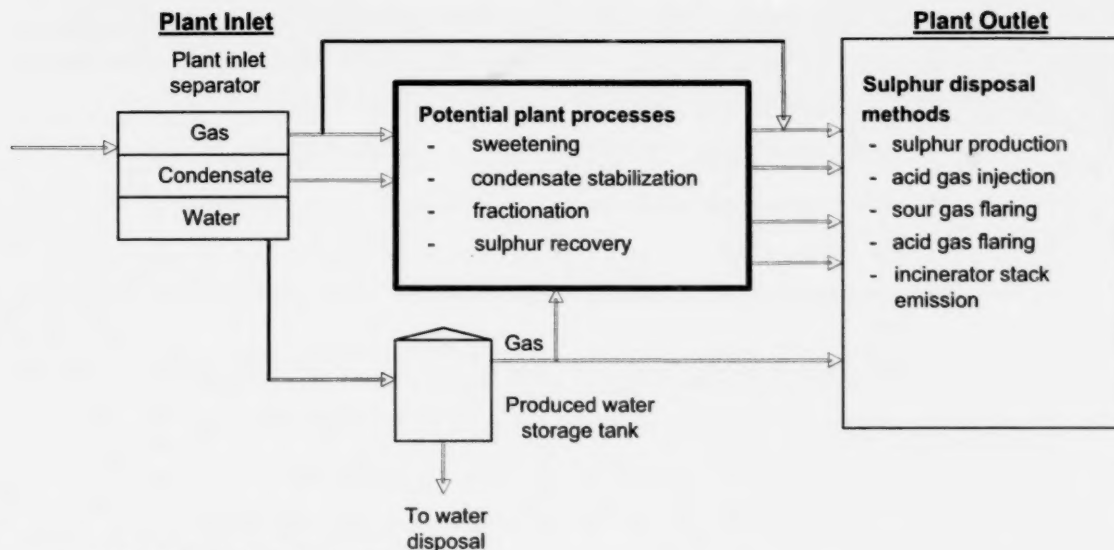
Figure 11.2. Sour gas plant process overview

As long as the gas and condensate entering a sour plant originate from a single pool, the H_2S concentration in the condensate will likely remain quite stable at the sampled conditions of pressure and temperature. Minor changes in pressure and temperature of the separator will only have a slight influence on the composition of the condensate. If major changes in pressure occur, such as due to installation of plant inlet compression and the resultant lowering of the inlet pressure, new samples must be taken and analyzed.

If the production to the plant occurs from two or more pools with different reservoir fluid compositions, the composition of the condensate will vary. In such cases, a vapour/liquid equilibrium correlation between the mole fraction of H_2S in the sour gas and the condensate can be used to estimate the mole fraction of H_2S in the condensate based on compositional analysis, computer process simulation, or stabilizer overhead volume and % H_2S to ensure acceptable accuracy for the S-30 report balance.

11.4.2.3 Determining Concentration of H₂S in Inlet Separator Water

The concentration of H_2S dissolved in the water that enters the plant inlet separator is a function of the separator pressure and temperature, as well as the concentration of H_2S in the sour separator gas. The amount of H_2S dissolved in the water can be predicted quite readily with correlations based on compositional analysis or computer process simulation.

Water can be metered ahead of the liquid level control valve on the water outlet line from the separator, or it can be gauged in a low-pressure produced water storage tank. The vapours from the tank are usually swept from the tank to flare with sweet gas. Some plants producing large amounts of sour water have installed a vapour recovery system, by which the tank vapours are conserved and treated in the processing plant.

If the total estimated sulphur content dissolved as H_2S in the sour water is less than 0.05 t/d (50 kg/d), the amount may be ignored in the balance determination.

It is recognized that a portion of the H_2S in the water of the plant inlet separator will remain in the water when the water is disposed of in a disposal well. This amount of H_2S is small and will depend on the water temperature in the produced water storage tank at atmospheric

pressure, the amount of agitation in the water, and whether sweet gas is used for sweeping the vapours from the tank to flare. The small amount of H_2S remaining in the water is difficult to estimate and therefore need not be included in the disposal accounting.

11.4.3 Calculation Procedure for Estimating the Plant Sulphur Inlet Mass per Day

1) The following streams must be accurately metered:

   a) sour gas out of the separator, Q, 10^3 m^3/d,

   b) sour condensate out of the separator, converted to gas equivalent volume, 10^3 m^3/d, and

   c) sour water out of the separator or into the storage tank, m^3/d, if required

2) The sulphur content in the sour gas out of the separator can be calculated:

   Sulphur equivalent in sour gas, t/d = (Q, 10^3m^3/d) (y) (1.35592)

   where y is the mole fraction of H_2S.

3) The condensate must be sampled and analyzed semiannually as a minimum frequency, in accordance with Section 8. When there are continuous gas analyzers and the H_2S content in the gas stream changes, the sulphur content in the condensate out of the separator can be calculated on the basis of the mole fraction of H_2S in the separator gas. The formula below should be used. Any alternative methods used must be supported by documentation that it is equivalent to the above method and made available to the ERCB upon request.

   x = y/K

   where: x = mole fraction of H_2S in the separator condensate

   y = mole fraction of H_2S in the sour gas in the plant inlet separator

   $K = A + (B-A) (T/66)^{1.2}$

   $A = -0.7034 (LOG_{10}(P))^3 + 9.1962 (LOG_{10}(P))^2 -39.58 LOG_{10}(P) +56.695$

   $B = -3.9694 (LOG_{10}(P))^3 +46.021(LOG_{10}(P))^2 -178.95 LOG_{10}(P) + 234.35$

   T = temperature of the sour gas in the plant inlet separator or metering temperature, °C

   P = pressure of the plant inlet separator, kPa(a)

   Sulphur equivalent in condensate, t/d = (Gas equiv. of condensate, 10^3m^3/d) (x) (1.35592)

   The range of applicability of the above method for determining x, the mole fraction H_2S in condensate, is between 700 to 9000 kPa(a) and 0 to 80°C.

4) The amount of H_2S dissolved in the water, z (mole fraction), in the plant inlet separator can be estimated by the following formula:[4]

   z = y/(4.53 – 7494.6/P + 758.4 (1.8 T + 32)/P + 4.65 y)

   where all terms are as defined above.

   Sulphur equivalent in water, t/d, = (1.31) (water production, m^3/d) (z) (1.35592)

---

[4] Froning, H. R. Jacoby, R. H., and Richards, W. L., "New K-Data Show Value of Water Wash," *Hydrocarbon and Petroleum Refiner*, April 1964, pp. 125-130.

The sum of the results of points 2, 3, and 4 for each sour gas inlet separator is the total sulphur inlet to the plant in t/d.

11.4.4 Calculation Procedure for Estimating Plant Sulphur Outlet Mass per Day

There are basically three different sulphur disposal schemes approved by the ERCB:

- sulphur recovery,
- acid gas flaring, and
- acid gas compression and injection.

Each of these schemes is treated separately as far as collecting the disposition data for the S-30 report is concerned. The plant inlet data are collected identically for the above different sulphur disposal schemes.

Potential sulphur disposal methods from sour gas plants are

- sulphur recovery,
- sour gas flaring or incineration,
- acid gas flaring or incineration,
- sulphur plant incinerator stack emissions, and
- sour gas flaring or incineration from the produced water storage tank (> 0.05 t/d).

The disposal of the sulphur by any of these methods must be accounted for. This requires measurement of flow rates and knowledge of concentrations of H_2S in the gas streams.

An important feature of the sulphur balance on the outlet side is the determination of the H_2S content of the acid gas out of the reflux drum. This gas stream is fully saturated with water vapour at the operating pressure and temperature of the reflux drum. Depending on what method is used in the determination of the H_2S content, the results could be on a dry basis or a wet basis. The operator must determine on which basis the analysis is determined.

The water content of the acid gas out of the reflux drum can be estimated by the procedure in Section 11.2.1.

Any H_2S determination and any complete analysis of the acid gas stream from the reflux drum presented on a dry basis must be normalized to a wet basis by the inclusion of the water vapour mole fraction. If the H_2S content in the acid gas is determined on a wet basis, the water vapour content is simply included as calculated above. In any case, the wet acid gas composition is to be used in the metering calculations of the acid gas stream at low pressure. This stream is then converted to a dry basis for reporting purposes.

11.4.4.1 Approved Sulphur Recovery Plants

The production of liquid sulphur must be determined by gauging the liquid sulphur level in sulphur production and storage pits or from weigh bills of shipments by truck or sulphur railroad tank cars, plus inventory changes in the pit.

Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flow rate is greater than $0.5 \ 10^3 \ m^3/d$ on a yearly basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter. A separate acid gas meter designed for the expected flow conditions and range

must be used to measure acid gas flared, either continuously or in emergencies, from gas sweetening systems regardless of volume.

The emissions from the sulphur plant emission stack must be monitored for flow rate and SO_2 concentrations.

The emissions from the vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations. These emissions must be reported as flared gas when this gas is not recycled or directed to the incinerator. If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur outlet of the plant but would still be a part of the sulphur inlet.

The sum of the sulphur contained in the above points must be the sulphur out of the plant. The difference between sulphur in and sulphur out of the plant must be no greater than ±5% if the actual inlet is ≥1 t/d or ±20% if the actual inlet is <1 t/d. The acid gas sent to the sulphur plant is to be reported as shrinkage (SHR), and acid gas flaring at the plant is to be reported as "FLARE" on the PRA.

11.4.4.2 Approved Acid Gas Flaring Plants

Plants approved for flaring of acid gas must meter the acid gas leaving the reflux drum of the sweetening process train. The meter calculation procedure must include the effect of the water vapour content in the acid gas at reflux drum pressure and temperature. It is recommended that the concentration of the H_2S content of the acid gas stream be checked at least once per week by Tutweiler by a person trained in the use of the technique and the calculation procedure to determine the H_2S concentration in the acid gas. A gas chromatograph may also be used for this analysis. Plants slipping CO_2 into the sales gas or receiving sour gas from different pools having different H_2S concentrations in the sour inlet gas may need to determine the H_2S concentration in the acid gas more often. A file must be set up to provide a record of the H_2S analysis determinations for inspection by the ERCB.

Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flow rate is greater than $0.5 \ 10^3 \ m^3/d$ on a yearly basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter.

The emissions from the vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations and must also be reported as flared gas if > 0.05 t/d. If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur out of the plant.

The sum of the sulphur contained in the above points must be the sulphur out of the plant. The difference between sulphur in and sulphur out of the plant must be no greater than ±5% if the actual inlet is ≥1 t/d or ±20% if the actual inlet is <1 t/d. The acid gas flared is to be reported on the PRA as shrinkage (SHR) and reported as "Flared" on the S30 report.

11.4.4.3 Approved Acid Gas Injection Plants

Plants approved for injection of acid gas into downhole injection wells must meter the acid gas leaving the reflux drum of the sweetening process train or at some point in the process piping at the plant site. If the gas is metered before the first stage of compression, the meter

calculation procedure must include the effect of the water vapour content in the acid gas at the reflux drum pressure and temperature. It is recommended that the concentration of the H_2S content of the acid gas stream be checked at least once per week by Tutweiler or gas chromatography by a person trained in the use of the technique and the calculation procedure to determine the H_2S concentration in the acid gas. Plants slipping CO_2 into the sales gas or receiving sour gas from different pools having different H_2S concentrations in the sour inlet gas may need to determine the H_2S concentration in the acid gas more often than once per week. A file must be set up to provide a record of the Tutweiler determinations for inspection by the ERCB.

Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flared gas flow rate is greater than $0.5 \ 10^3 \ m^3/d$ on a yearly basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter. A separate acid gas meter designed for the expected flow conditions and range must be used to measure acid gas flared, either continuously or in emergencies, from gas sweetening systems regardless of volume.

The emissions from the vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations and must also be reported as flared gas if > 0.05 t/d. If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur out of the plant.

The sum of the sulphur contained in the above points must be the sulphur out of the plant. The difference between sulphur in and sulphur out of the plant must be no greater than ±5% if the actual inlet is ≥1 t/d or ±20% if the actual inlet is <1 t/d. The acid gas injected is to be reported as a disposition (DISP) to the injection facility, and acid gas flaring at the plant is to be reported as "FLARE" on the PRA.

11.5   Production Data Verification and Audit Trail

The field data, records, any calculations or estimations, and EFM records relating to ERCB-required production data submitted to the PRA and the ERCB S-30 reports must be kept for inspection upon request. The records verification and audit trails must be in accordance with the following:

- Proving/calibration records: any records and documentation produced in the proving/calibration of meters and calibration of the prover and all peripheral devices (if the prover and peripheral devices are owned and operated by the licensee)

- Delivery and receipt records: any records and documentation produced in the determination of delivery or receipt volumes/tonnage

- Estimation records: any records and documentation related to the estimation of reported volume/tonnage, including estimation methodology, record of event, and approvals

- Pit gauging records: any records and documentation produced in the determination of reported volume/tonnage

- Volume/tonnage loss records: any records and documentation for volumes lost due to incidents such as spills and fires

- EFM: any records and documentation (electronic, magnetic, or paper form) produced in the determination of measured volume/tonnage in accordance with the EFM requirements in Section 4.3.3.2 for gas and Section 6.8 for liquids.

11.6 How to Complete the S-30 Monthly Gas Processing Plant Sulphur Balance Report

The S-30 Monthly Gas Processing Plant Sulphur Balance Report below must be submitted to the ERCB using the electronic Digital Data Submission (DDS) system under "Submit Monthly Sulphur Balance Reporting," according to the instructions that follow the form.

ERCB *Energy Resources Conservation Board*

Monthly Gas Processing Plant Sulphur Balance Report (S-30)
Gas Volumes in $10^3 \, m^3$ at 101.325 kPa and 15°C
Sulphur Measurements in Tonnes (t)

Plant Name _____

Operator _____

Location

LSD	SEC	TWP	RGE		M
					W

Monthly Measurement Diff (%) _____

Emission Monitor Down (h) _____

PLANT CODE	YEAR	MONTH

Day	Sulphur In			Sulphur Out								
	Plant Feedstock			Sulphur Production	Sulphur Stack Emission	Flared Gas			Injected / Other			Total
	Approved Max. ($10^3 \, m^3$)		Approved Max. (t)									
	Actual $10^3 \, m^3$	Actual % H_2S	Actual t	t	t	$10^3 \, m^3$	% H_2S	t	$10^3 \, m^3$	% H_2S	t	t
1												
2												
3												
4												
5												
6												
7												
8												
9												
10												
11												
12												
13												
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21												
22												
23												
24												
25												
26												
27												
28												
29												
30												
31												
Month Total												
Qtr Total												

Sulphur Recovery Efficiency	Minimum Approved %	Actual Monthly %	Cumulative Quarterly %

Contact Name _____   Telephone _____

Signature _____   Date _____

S - 30 - 2006-05

Energy Resources Conservation Board 640 - 5 Avenue SW Calgary, Alberta Canada T2P 3G4

How to Complete the S-30 Monthly Gas Processing Plant Sulphur Balance Report

Identification

Plant Name	Enter the name of the gas processing plant as given on your gas processing facility approval or from *ST50: Gas Processing Plants in Alberta* (typically licensee name and the first field name).
Operator	Enter the operator name.
Location	Enter the legal description of the plant as shown on the approval.
Monthly Measurement Difference (%)	This field will be calculated using the following formula:

Monthly measurement difference (%) = [Monthly sulphur in (t) − Monthly sulphur out total (t)] ÷ Monthly sulphur in (t)

Emission Monitor Down (h)	Enter the number of hours the continuous stack emission monitor (CSEM) was not in service during the reporting month.
Plant Code	Enter the gas processing plant facility code. The field size is 4 numeric characters.
	• The facility code must exist on the PRA.
Year and Month	Enter the 4-digit year and the 2-digit number of the month being reported. The field size is 6 numeric characters.

Reporting Data (General)

You must file an S-30 report if you are an operator of a gas processing plant in which raw gas is processed for the removal of H_2S.

Report gas volumes in 10^3 m^3 at 101.325 kPa pressure and 15°C.

Report sulphur in tonnes (t) to 1 decimal place, with the option to report to 2 decimal places if required to meet measurement difference requirements.

Day	Enter the data on the row of the corresponding day of the month.

Sulphur In

Plant Feedstock— **Approved Maximum (10³ m³)**	Enter the approved maximum volume of plant feedstock (raw gas plus gas equivalent of inlet condensate), as stated in the approval or the most recent application approved (see Schedule 2, *Directive 056: Energy Development Applications and Schedules*) in 10^3 m³ to 1 decimal place.
Plant Feedstock— **Approved Maximum (t)**	Enter the approved maximum of inlet sulphur in tonnes per day to 1 decimal place, as stated in the approval or the most recent application approved (see Schedule 2, *Directive 056*). Enter the daily values for sulphur inlet and outlet as detailed below in the row alongside each appropriate date.
Plant Feedstock **(10³ m³)**	Enter the actual volume of plant feedstock (raw gas plus gas equivalent of inlet condensate) received by the plant on this day in 10^3 m³ to 1 decimal place (see Section 8.3). This must not be entered as a daily average volume for the entire month.
Plant Feedstock **(% H₂S)**	Enter the daily percentage of H_2S contained in the recombined plant feedstock (raw gas and condensate and/or water) on this day to 1 decimal place; if less than 0.10%, enter the percentage to 4 decimal places.
Plant Feedstock (t)	This field will be calculated using the following formula: Sulphur (t) = Plant feedstock volume (10^3 m³) × Recombined H_2S% × 1.35592 (Conversion factor) ÷ 100

Sulphur Out

Sulphur Production (t)	Enter the daily sulphur produced in tonnes to 1 decimal place, with the option to report to 2 decimal places if required to meet measurement difference requirements.
Sulphur Stack Emission (t)	Enter the daily sulphur emissions in tonnes to 1 decimal place, with the option to report to 2 decimal places if required to meet measurement difference requirements as recorded by the continuous stack emission monitor (CSEM).
Flared Gas (10³ m³)	Enter the amount of gas flared from the plant in 10^3 m³ to 1 decimal place. • Include all sour gas flared, • In this column, also report acid gas flared from plants that do not recover sulphur.
Flared Gas (% H₂S)	Enter the daily percentage of H_2S contained in the flared gas to 1 decimal place.

Flared Gas (t)	This field will be calculated using the following formula:
	Sulphur (t) = Flared gas volume (10^3 m^3) × Flared gas H$_2$S% × 1.35592 (Conversion factor) +100
Injected/Other (10^3 m^3)	Enter the daily volume of sulphur injected to subsurface formations or disposed of in any manner other than described above (e.g., nonregenerative sweetening) in 10^3 m^3 to 1 decimal place.
	• Include all sour gas injected.
	• In this column, also report acid gas injected from plants that do not recover sulphur.
	• "Other" disposition, if used, must be identified and separately quantified (monthly total) as part of the S-30 submission.
Injected/Other (% H$_2$S)	Enter the daily percentage of H$_2$S contained in the injected gas to 1 decimal place.
Injected/Other (t)	This field will be calculated using the following formula:
	Sulphur (t) = Injected gas volume (10^3 m^3) × Injected gas H$_2$S% × 1.35592 (Conversion factor) +100
Total (t)	This field will be calculated using the following formula:
	Total daily tonnage = Sulphur production (t) + Sulphur stack emission (t) + Flared gas (t) + Injected/others (t)
Month Total	This field will be calculated automatically.
Quarterly Total	This field will be calculated automatically.
	• The ERCB defines the quarters as January-March; April-June; July-September; and October-December. Thus, for example, if you are filing for February, enter the monthly total for January plus February; if filing for March, enter the monthly total for January plus February plus March; if filing for April, enter the monthly total only for April.

Sulphur Recovery Efficiency

Complete this section only if the plant recovers elemental sulphur or injects acid gas.

Minimum Approved %	Enter the approved minimum quarterly sulphur recovery efficiency as stated in the approval or from *ST50: Gas Processing Plants in Alberta*.
Actual Monthly %	This field will be calculated using the following formula:

Actual monthly sulphur recovery efficiency =

$$\frac{(\text{Sulphur production (t)} + \text{Sulphur injected}) \times 100}{\text{Sulphur production (t)} + \text{Emissions (t)} + \text{Flared (t)} + \text{Sulphur injected}}$$

Cumulative Quarterly %	This field will be calculated automatically.
Contact	Enter the full name of the person who accepts responsibility and to whom inquiries regarding this report should be directed.
	Sign the statement.
	Enter the person's business telephone number, including area code. (Please print.)
	Enter the date this S-30 report is completed.
When to File	Your monthly S-30 report must be submitted to the ERCB Operations Group via DDS by 4:30 p.m. on the 18th day of the month following the month being reported. If the 18th day is not a business day, you must submit on the first business day after the 18th.

12 Heavy Oil Measurement

12.1 Introduction

This section presents the requirements and exceptions for heavy oil and crude bitumen facilities. This section does not cover crude bitumen production through mining.

The term "heavy oil" in this section includes crude bitumen (other than that produced through mining or in situ processes). See Glossary for definition of heavy oil and crude bitumen.

There are two general categories for the production of heavy oil: primary/secondary (includes cold production and water flood techniques) and in situ thermal and solvent oil sands schemes.

Blending of heavy oil with lighter hydrocarbon liquids may occur. The lighter hydrocarbon liquids, often called diluent or condensate, are used to reduce the viscosity of the heavy oil and make it easier to process and/or transport.

12.2 General Measurement Requirements

In all cases, a licensee must measure produced heavy oil volumes, unless otherwise stated in this directive. The ERCB will consider a heavy oil measurement system to be in compliance if the fuel measurement requirements in Sections 1.7 and 4.2, the calibration and proving requirements in Section 2, and measurement requirements in Section 6.3 and in this section are met.

12.2.1 Temperature Correction Requirements

See Section 6.3.2.1: Temperature Correction Requirements

Exception

If testing heavy oil by test tanks, temperature correction is not required. However, the operator must be aware that if the temperatures are elevated above standard conditions, the proration factors will be biased by the amount of the temperature correction (approximately 0.08 per cent per degree Celsius [%/°C] at a density of 920 kg/m^3). See section 3.1.2 for required amendments.

12.2.2 Pressure Correction Requirements

See Section 6.3.2.2: Pressure Correction Requirements.

12.2.3 EFM Requirements

See Section 6.8: Electronic Flow Measurement for Oil Systems.

12.2.4 Diluent/Condensate Receipts and Blending

Blending occurs when two oils of dissimilar properties are mixed. This mixing result in volumetric discrepancies from the ideal combination, which would yield a volume that would be the sum of the two products. The discrepancy is usually shrinkage, which is the result of smaller molecules of the lighter hydrocarbon filling in the voids or spaces between larger molecules of the heavier hydrocarbon. The result is a combined liquid volume that is less

than the sum of the two original volumes. This shrinkage must be determined and properly applied to volumes making up the liquid to ensure proper allocation and reporting.

The diluent used for blending purposes is often a royalty paid product and measured at the receipt point. This means that the diluent injected is netted off of the sales products prior to determination of produced or sales gas, oil, or bitumen.

12.2.4.1   Hydrocarbon Blending and Flashing Shrinkage

Generally, heavy oil is not significantly affected by shrinkage caused by flashing of light ends. However, there is always potential for shrinkage depending on the actual operating pressure at which the wells are produced. Operators should evaluate each facility based on its operating characteristics.

Blending of diluent with heavy oil can occur at any point in the production process, such as in the flow line from the well, at the inlet separator, at the treater, or at the storage tank. If diluent is blended with the oil prior to the treater, diluent flashing (vapourization) may also occur. Operators must determine and account for both the blending and the flashing shrinkages using engineering estimates and report them on the PRA. The blending shrinkage must be reported as an "SHR" disposition of the facility, and the flashing shrinkage must be reported as a GEV (ABGE) of diluent receipt (REC) into the facility. When reporting the shrinkage, either flashing or blending shrinkages must be applied to the diluent volume, and the heavy oil volume must not be reduced by the shrinkage.

Blending shrinkage calculation methods can be found in API MPMS, Chapter 12, Section 3. Equivalent industry-accepted methods may also be used. Diluent flashing shrinkage can be estimated using computer simulation programs based on the process conditions with the composition and density of the diluent and heavy oil. See Appendix 8 for an example.

12.2.5   Water Measurement

Operators must have in place proper procedures to measure produced water, water receipts, and water injection/disposal to ensure that the information used for reporting is accurate. See Section 1 for measurement accuracy requirements.

As sand may constitute a significant percentage in the S&W content of heavy oil production, it is expected that operators will follow Section 6.4 requirements for sampling and analysis to quantify these volumes. The sand volume is to be included as part of the S&W determination and reported as water.

12.2.6   Well Proration Testing

For heavy oil well testing, operators must follow the testing frequency requirements in Section 6.4.4. For additional heavy oil well testing requirements, see Sections 12.3.4.

12.2.7   Heavy Oil Receipt, Delivery, or Sales

This includes the delivery or receipt of heavy oil (diluted or non-diluted), diluent, or other hydrocarbon products. For the single point measurement uncertainty of these measurement points see Section 12.3.3.

12.3 Primary and Secondary Production (Non In Situ Thermal and Solvent Oil Sands Schemes)

Well effluent produced from heavy oil wells using natural or other drive mechanisms, such as water-flood wells, flowing wells, or wells with conventional lifting technologies (pump jacks, progressive cavity pumps, submersible pumps, etc.), is considered primary and secondary production. The well effluent could be flow lined to field test satellites or group production facilities, or the fluids could be produced to tanks at surface and trucked out to a treatment facility. These production wells are subject to the same measurement and reporting requirements as conventional oil wells (see Section 6), unless otherwise noted in this section.

12.3.1 Battery Types

Battery types are the same as in Section 6.2.2, with the following exceptions:

12.3.1.1 Single-Well Batteries—PRA Subtype 331 (Disposition = Production)

Oil and water volumes trucked from lease production tanks must be used to calculate the well production reported on the PRA. Where a well is producing as a single-well battery (SWB) to a lease tank, the lease tanks can be considered as part of the reservoir and the inventory in the tanks is not reported. This procedure is referred to as the disposition equals production (disposition = production) accounting method. Oil and water production are only reported when the fluids are trucked from the lease tank to another location.

If a well is on a restricted gas production order (gas allowables), the disposition = production method may not be appropriate and the reporting of oil and gas production should be done monthly based on inventory change. If there is receipt(s) into the single-well battery, monthly production must be reported based on receipt and inventory volumes and the disposition equals production methodology does not apply.

When using the disposition = production accounting method, it is correct to show hours on production and no production volume if a shipment was not made from a lease tank of a producing well during the reporting period. Conversely, produced fluid removed from a lease tank during a month that a well is shut in is to be shown on the reports as with zero hours of production. Suspended and abandoned heavy oil batteries can report dispositions (and production) up to six months after they become inactive.

12.3.1.2 Paper Batteries—PRA Subtype 343

Paper batteries are developed to reduce the number of reporting entities for submission to the PRA. Paper batteries are made up of multiple single heavy oil well batteries (excluding multiwell group batteries) that are grouped for reporting purposes into one larger battery. This allows operators to report production and dispositions from a number of individual wells in one battery, thereby reducing the administrative burden. Paper batteries are treated as multiwell group batteries even though the single wells are not on a common production site. On the PRA, a paper battery is to be reported as a subtype 343: Crude Bitumen/Heavy Oil Administrative Grouping.

It is acceptable to move fluids between locations within the same paper battery. These volume movements are not reported to the PRA but must be managed in a field data capture system.

Wells in a paper battery are eligible for reporting using the disposition = production methodology.

As paper batteries are for administrative ease, they must comply with the following criteria:

- Wells within the paper battery must have common equity and royalty.
- No venting or fuel can be reported at the paper battery level. Venting and fuel activity must be reported at the location where it physically occurred, i.e., at the well site.
- Wells within a paper battery must be within a geographic area no larger than 6 contiguous townships.
- The operator must use one of the SWB locations within the paper battery for reporting.

12.3.1.3 Multiwell Group Batteries—PRA Subtype 341

Each well must have its own separation and measurement equipment similar to a single well battery. The disposition = production accounting method may be used for each well when appropriate.

All equipment for the wells in the battery must share a common surface location.

Inventory at the group battery is to be reported monthly to the PRA.

12.3.1.4 Multiwell Proration Batteries—PRA Subtype 342

All well production is commingled prior to the total battery oil/emulsion being separated from the gas and measured. Individual monthly well oil and water production is estimated based on periodic well tests and corrected to the actual monthly volume through the use of a proration factor (see Section 6.4.4).

Inventory at the proration battery is to be reported monthly to the PRA.

Wells included in a proration battery are not eligible for the disposition = production methodology.

12.3.2 Gas Measurement and Reporting

Any single stream of produced gas, flared gas, or vented gas volume exceeding $2.0 \ 10^3$ m^3/day must be measured. If the annual average fuel gas usage exceeds $0.5 \ 10^3 \ m^3$/day on a per site basis, the fuel gas must be measured.

For single-well batteries, vented or flared gas at a well site may be reported at the battery level. For a multiwell group, proration, or paper batteries, individual well flared or vented gas must be reported at the well level unless the gas is collected at a common point and then flared or vented. Then it can be reported at the battery level.

If gas measurement is required, the single point measurement uncertainty is ±3%.

12.3.2.1 Single-Well Batteries

For initial well startup, in the absence of suitable reservoir information, monthly tests must be conducted to determine the GOR factor or the hourly rate if gas volumes are not dependent upon oil production volumes (see Section 4.3.5) for 6 months or until gas production stabilizes and measurement is required if over $2.0 \ 10^3 \ m^3$/day. If total gas production per well does not exceed $2.0 \ 10^3 \ m^3$/day, the producer may use GOR or hourly rate testing to determine gas volumes. In addition to the measurement of volume, the source of the produced

gas must be assessed by analysis to determine whether the gas is solution gas or cap gas. This is especially important if gas lease ownership differs from bitumen lease ownership.

A test of the well (approximately 24 hours) may be conducted such that all the applicable gas and oil volumes produced during the test are measured (including fuel, flared, and vented gas; see Figure 12.1). The gas volume is to be divided by the oil volume to result in the GOR factor or divided by the test hours to obtain an hourly rate. For single-well oil batteries the oil volume used in the determination of the GOR factor can be based upon the monthly total oil production (monthly total volume / hours produced in month x test duration).
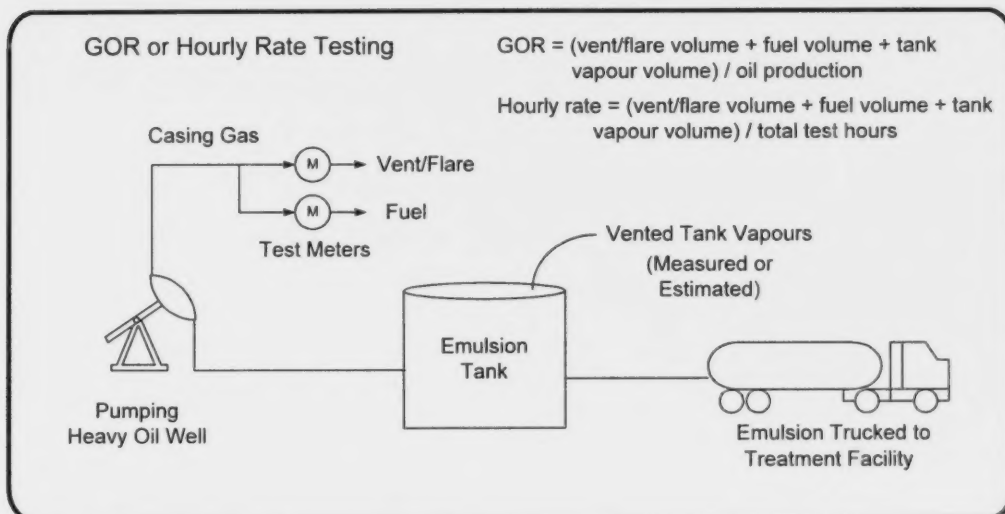
Figure 12.1. GOR or hourly rate testing schematic

GOR or hourly rate test frequencies are detailed in the table below.

Table 12.1.   GOR or hourly rate test frequency requirements

Gas rate (10^3 m³/d)	Test frequency
≤0.1	Once every 3 years
>0.1 but ≤1.0	Annually
>1.0 but <2.0	Semi-annually

12.3.2.2   Multiwell Batteries

If gas disposition is measured at a multiwell battery and is sold or used at a point farther on, well gas production volumes may be tested with measurement if over $2.0 \ 10^3$ m³/day or estimated using well GOR or hourly rate if not more than $2.0 \ 10^3$ m³/day (see Section 12.3.2.1).

For multiwell group or proration batteries, if associated gas is flow lined to a central facility or collection point and gas production rates at the point of collection or emission is less than $2.0 \ 10^3$ m³/day, a group or facility GOR may be determined (see Table 12.1), but gas production reporting must be at the individual well level. Initial and updated factors may be determined by any of the applicable tests or procedures described in Section 4.3.5.

12.3.3 Oil and Water Deliveries to a Treatment Facility

Liquid volumes are trucked from single-well batteries, multiwell batteries, and wells within paper batteries to a custom treater or battery. These facilities produce oil, water, and sand for disposal from the trucked-in fluids. Typically, received products are measured using weigh scales or inlet meters but tank gauging could also be used (see Section 10.3). The oil and water densities from every well must be updated in accordance with Section 10.4.4 and the S&W of the emulsion delivered to these facilities must be determined on a per-load basis (see Sections 6.4.2 and 6.4.3).

The single point measurement uncertainty of liquid products received into the cleaning plant is ±1%, excluding the effect of S&W and density determination. The single point measurement uncertainty of sales oil delivery point measurement from the treatment facility is ±0.5%.

12.3.4 Well Test Measurement with Tank Gauging or Metering

Wells in primary/secondary production of heavy oil must be tested at the frequency stated in Section 6.4.4. The tests must be conducted in a consistent manner throughout the month and a test must be conducted when there is a change in well parameters (pump speed, work-over, reactivation, flush-by, etc.) as soon as possible.

A water cut must be determined for each test. See Sections 6.4.2 and 6.4.3.

Temperature correction is not required for well tests (see Section 12.2.1).

Tests may be conducted by using a single isolated tank at the well site. The table below may be used by the licensee to determine the height-to-diameter ratio requirements in accordance with Section 6.3.1.2a, with the exception that the accuracy coefficient in this table is a suggested minimum for test fluid volumes.

Table 12.2. Accuracy coefficients for various measurement types for test tanks

Level measurement technique	Accuracy coefficient "a"	Maximum level reporting resolution (mm)
Gauge board	1.6	25
Manual dip of the tank	0.4	10
Electronic (e.g., radar)	0.4	10

The accuracy coefficient "a" can be used in the following equation:

$$V \geq a \times d^2 \quad \text{or} \quad d \leq (V/a)^{0.5}$$

where:

V = test fluid volume in m^3
a = accuracy coefficient
d = tank diameter in metres

On the tank being used for testing, the gauge board float, linkage, and scale must be in good condition. The gauge board markings must be no further than 60 mm apart. For gauge measurement on test tanks, one reading of the gauge board is acceptable for the start and end of the test. Where safe work conditions permit, gauge boards should be read at eye level.

If the well emulsion is to be tested using a meter, the meter must be sized to operate within 20 to 90% of its flow range and installed and operated in accordance with Sections 2.6.1 and 6.3.1.1, except no temperature measurement is required.

The single point measurement uncertainty of the test emulsion meter is ± 2%, excluding the effect of S&W determination.

12.4 In Situ Thermal and Solvent Oil Sands Schemes

This section will be added once further research and consultation are completed.

12.5 Uncertainty Summary

Table 12.3. Summary of single point measurement uncertainty

Measurement type	Single point measurement uncertainty
Gas in primary/secondary production or injection	±3%
Emulsion test using metering	±2%, excluding the effect of S&W determination
Primary/secondary production emulsion into treatment facilities	±1%, excluding the effect of density and S&W determination
Clean oil sales	±0.5%

13 Condensate and High Vapour Pressure Liquid Measurement and Reporting

This section presents the requirements and exceptions for condensate measurement and reporting associated with gas well production.

Condensate is a generic term used to describe various types of hydrocarbon liquid products. This section is intended to clarify the various definitions of condensate and address the measurement and reporting requirements. See Glossary in Appendix 3 for definition of high vapour pressure liquids, NGL, LPG, ethane, propane, butane, and pentane plus.

13.1 Definitions

Field condensate: For Crown Royalty purposes, field condensate (in some documents, this is simply referred to as condensate) is defined as "products obtained from natural gas or solution gas before they are delivered to a gathering system." Typically, Field Condensate is a hydrocarbon liquid separated from raw production at a well or a group measurement point, stabilized in a tank, and sold or otherwise disposed of without further processing before entering a gas gathering system. See *Natural Gas Royalty Regulation 2002*, AR 220/2002 (*NGRR*), under the *Mines and Minerals Act*.

Condensate, separator liquids: Separator liquids are a mixture of hydrocarbon components that remain in a liquid state under the equilibrium pressure and temperature (P&T) conditions established in a two- or three-phase separator. The composition and physical properties of separator liquids are highly variable and are a function of separator inlet fluid composition and separator P&T conditions.

Condensate, stock tank liquids: Stock tank liquids are a mixture of hydrocarbon components that remain in a liquid state under the equilibrium P&T conditions in a low-pressure vessel or atmospheric-pressure tank. The composition and physical properties of stock tank fluids are a function of the composition of the high-pressure fluid from which they were derived.

Condensate, stabilized liquids: Stabilized liquids are a mixture of hydrocarbon components that remain in the liquid state following single-stage flash evaporation. Stabilized condensate is expected to have low concentrations of butane minus (C4-) components. The composition and physical properties of the stabilized condensate is a function of the equilibrium P&T conditions of the stabilizer and the composition of the stabilizer feed from which it was derived.

13.2 General Measurement and Reporting Requirements

13.2.1 Measurement Requirements

For measurement of condensate and high vapour pressure liquids at all accounting locations within the upstream oil and gas facilities, the ERCB will consider the measurement system to be in compliance if the measurement requirements in Sections 1.7.2 and 4.2, the calibration and proving requirements in Section 2, the design and installation of liquid (oil) measurement requirements in Section 6.3, the sampling and analysis requirements in Section 8, and the trucked liquid requirements in Section 10 are met.

13.2.2   Reporting Requirements

Hydrocarbon liquid production can be reported as a gas or liquid or both, depending on how it is disposed (see *Directive 007: Volumetric and Infrastructure Requirements* (December 2007)). The general rules are as follows for PRA reporting:

- Hydrocarbon liquids produced from a gas well producing oil must be reported as a liquid oil volume at the well level, regardless of whether they are trucked or recombined. The *Petroleum Royalty Regulation 2008,* AR 248/90 (PRR), under the *Mines and Minerals Act* applies for Crown royalty.

- Hydrocarbon liquids received from wells designated as VGWL OIL that are separated at group separation points at batteries or gathering systems and directed to a tank for disposition are considered OIL and must be reported as a liquid volume.

- Hydrocarbon liquids produced and separated from a gas well or at the group measurement points of multiwell gas proration or effluent proration batteries producing condensate, measured, and recombined with the gas or trucked out for further processing must be converted to and reported as gas production (see Cases 1 and 5 below). The *NGRR* applies for Crown royalty.

- Hydrocarbon liquids produced and separated from a gas well producing condensate or at the group measurement points of multiwell gas proration or effluent proration batteries, stored in a tank, and trucked for sales are considered field condensate and must be reported as a liquid condensate volume at the well level (see Cases 2 and 3 below). The *NGRR* applies for Crown royalty.

- Hydrocarbon liquids separated from a gas gathering system at group separation points and stored in a tank for disposition are considered pentanes plus (C5+) and must be reported as a liquid volume (see Case 4 below). The *NGRR* applies for Crown royalty.

- Hydrocarbon liquids recovered from processing at a gas processing plant are considered by-products of processing and must be reported as a liquid volume. The *NGRR* applies for Crown royalty.

- Load condensate (or oil or water) received, injected, recovered, separated, and stored in a tank at a measured gas well is to report load fluids, in accordance with *Directive 007* on-line supplement Appendix 8 reporting procedures.

- Load condensate received from an outside source at any gas well, injected, recovered, separated, recombined with production, and sent to a gas plant must report condensate receipt at the battery level and report disposition of C5+ with From/To of "ABGE" to balance the condensate receipt to avoid double paying royalty.

- Load LPG, NGL, C3, or C4 received from an outside source at any gas well, injected, recovered, separated, recombined with production, and sent to a gas plant must report hydrocarbon receipt at the battery level and report disposition of the hydrocarbon with From/To of "ABGE" to balance the hydrocarbon receipt to avoid double paying royalty.

- Load oil received from an outside source at any gas well, injected, recovered, recombined with production, and sent to a gas plant must report oil receipt at the battery level and report disposition of the oil.

13.3 Reporting Scenarios

Gas Wells Producing Condensate

Condensate is associated with gas well production within gas facilities. If there is condensate, the total battery liquid volume must be measured.

Case 1

Gas, condensate, and water being wet metered and tested or proration tested only, or separated from other well effluent, metered, commingled with gas (single well or multiwell group batteries) and sent to a gas plant for further processing must be converted to its GEV and added to the gas production for reporting purposes. The *NGRR* applies to this volume as part of the total production and disposition at the gas plant for Crown royalty.
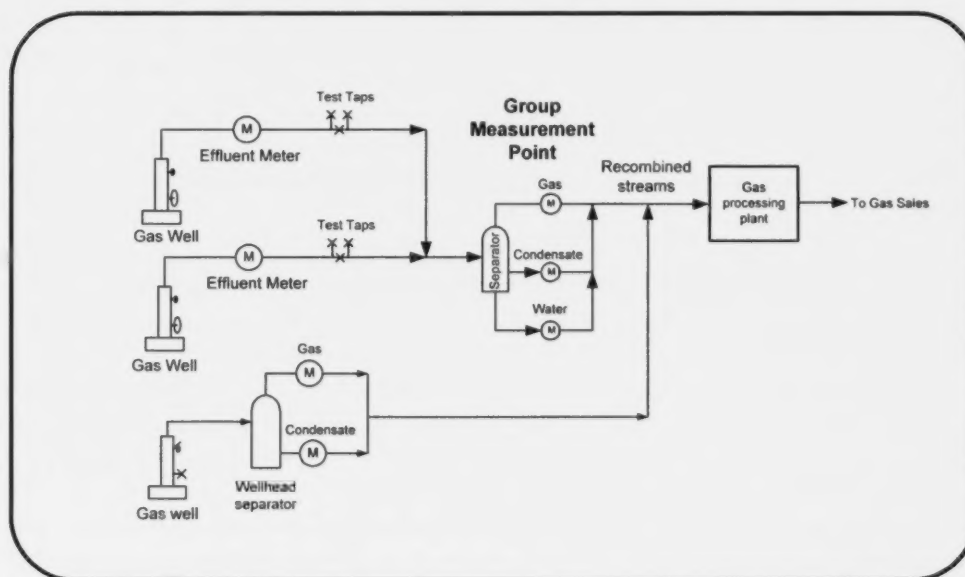
Case 2

Field condensate must be reported as a liquid production at the well if separated from well effluent, measured, and disposed of without further processing and before being delivered to a gas gathering system. The *NGRR* applies to this volume for Crown royalty.
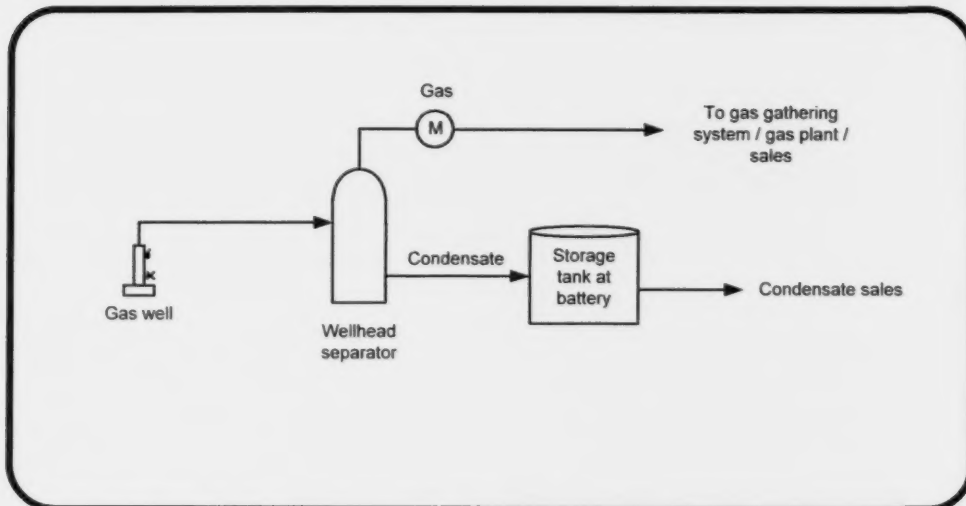
Case 3

Field condensate must be reported as a liquid production at the well level if separated from multiwell gas proration or effluent proration batteries, measured, and disposed of from the group separator without further processing and before being delivered to a gas gathering system. The *NGRR* applies to this volume for Crown royalty.
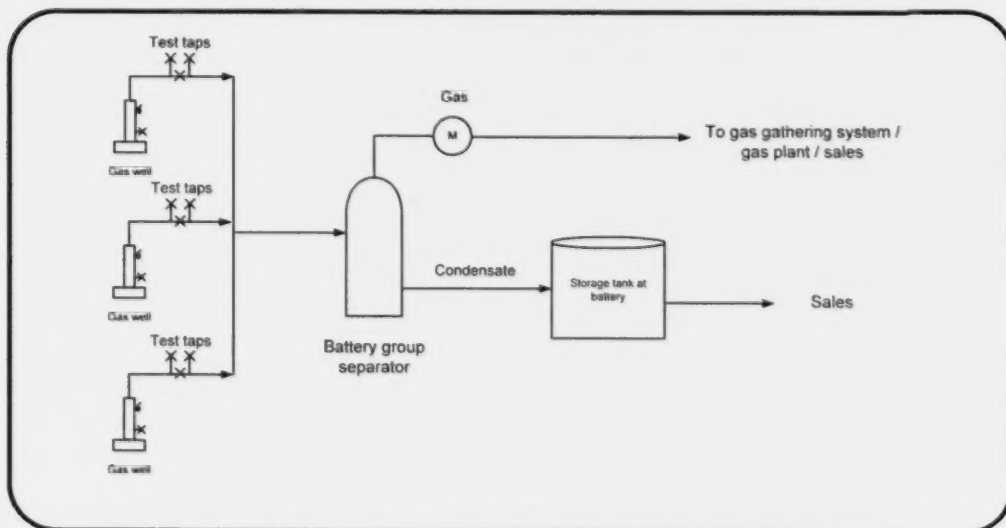
Case 4

Hydrocarbon liquid from one or more batteries recovered as a result of gas compression at a gas gathering system or a gas group battery and disposed of without further processing must be reported as pentanes plus liquid at the gas gathering system or gas group battery. The *NGRR* applies to this volume for Crown royalty.
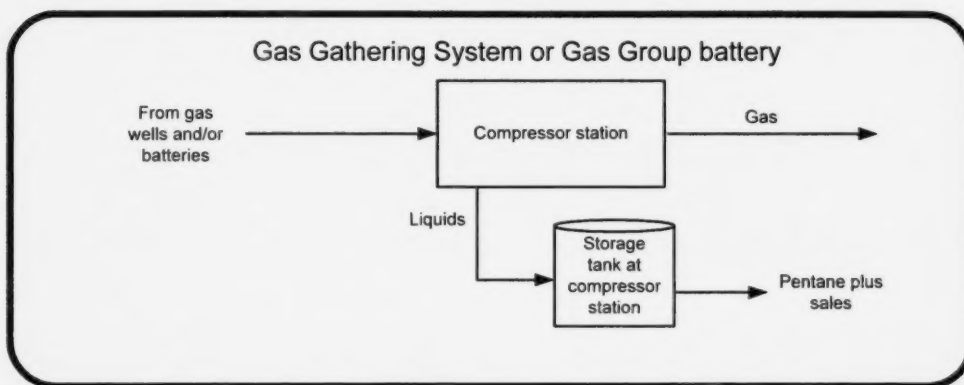
Case 5

Condensate separated from well effluent, measured, and trucked to a gas plant for processing must be converted to its GEV and added to the gas production for reporting purposes. The *NGRR* applies to this volume as part of the total production and sale at the gas plant. If there is no processing before sales, see Cases 2 and 3.
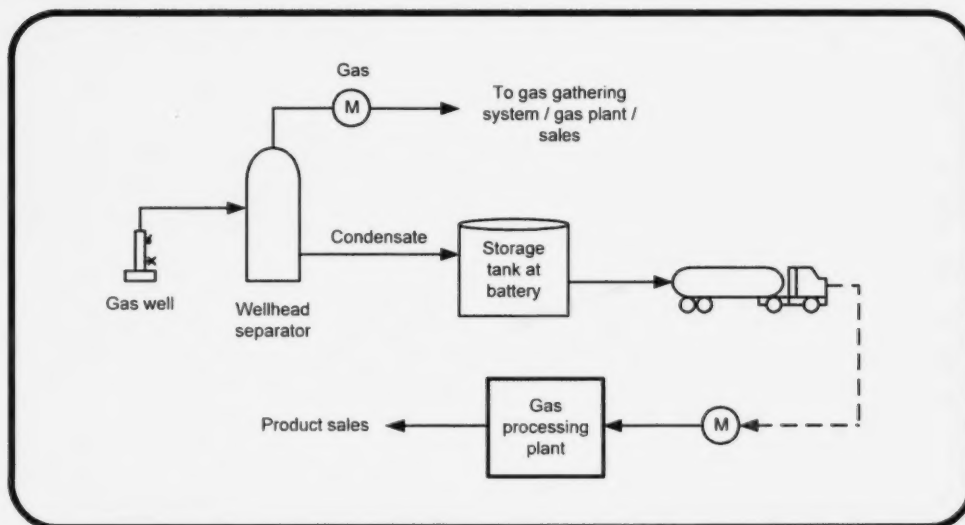
Appendix 1   Sections Under Development

Liquid Measurement

Water Measurement

Gas Lift Systems

Downhole Gas/Liquid Separation and Injection Systems

Appendix 2   ERCB Documents Replaced Fully or Partially by *Directive 017: Measurement Requirements for Upstream Oil and Gas Operations*

Total Replacement (rescinded on date *Directive 017* becomes effective):

Informational Letter (IL) 86-03: Automated Measurement Systems
IL 87-1: Compressibility Factors Used in Gas Volume Calculations and Physical Property Data for Natural Gases
IL 89-16: Guidelines for Automated Measurement System Applications, ERCB Guide G-34 Comparative Chart Data
IL 90-6: Measurement Guidelines—Trucked Oil Production
IL 91-9: Exemption from Gas Measurement Crude Oil/Bitumen Wells
IL 92-8: Crude Oil Pipeline Truck Terminal Measurement Guidelines
IL 93-1 Gas Density Measurement Frequency—Orifice Meter
IL 93-10 Revised Measurement and Accounting Procedures for Southeastern Alberta Shallow Gas Wells
IL 94-7: Coriolis Force Flowmeters
Interim Directive (ID) 90-2: Gas Meter Calibration
ID 91-3: Clarification (May 2001): Heavy Oil/Oil Sands Operations
ID 94-1: Measurement of Oil, Gas & Water Production
Directive 049: Gas Density Measurement Frequency

Partial Replacement:

For sections in ILs and IDs that are no longer effective, see the documents on the ERCB Web site www.ercb.ca:
ID 91-3: Heavy Oil/Oil Sands Operations

Directive 004: Determination of Water Production at Gas Wells
Directive 007: Production Accounting Handbook
Guide 34: Guidelines for Automated Measurement System Applications—Note that for those systems approved under *Guide 34*, the guide is still in effect.
Directive 046: Production Audit Handbook
Directive 060: Upstream Petroleum Industry Flaring Guide
Directive 063: Oilfield Waste Management Facility Inspection Manual
Directive 064: Facility Inspection Manual

Appendix 3 Glossary

The definitions that follow are for the purposes of this directive only.

Absolute Density of Gas - The gas's mass per unit volume at a specific pressure and temperature. Absolute densities are generally expressed in kg/m^3 at 101.325 kPa(a) and 15°C.

Absolute Density of Liquid - The liquid's mass per unit volume at a specific pressure and temperature. Absolute density of a liquid is generally expressed as kg/m^3 at 101.325 kPa(a) and 15°C.

Accuracy - The ability of a measuring instrument to indicate values closely approximating the true value of the quantity measured.

Acid Gas - Gas separated in the treating of solution or nonassociated gas that contains hydrogen sulphide (H_2S), totally reduced sulphur compounds, and/or carbon dioxide (CO_2).

Allowable - Used in connection with a well, means the amount of oil or gas a well is permitted to produce, in accordance with an order of the ERCB Board for this purpose, after application of any applicable penalty factor.

Bias - Any influence on a result that produces an incorrect approximation of the true value of the variable being measured. Bias is the result of a predictable systematic error.

Battery - A system or arrangement of tanks or other surface equipment receiving the effluents of one or more wells prior to delivery to market or other disposition, and may include equipment or devices for separating of the effluents into oil, gas, or water and for measurement.

Block - An area or part of a pool consisting of drilling spacing units grouped for the purpose of administering a common, aggregate production allowable.

Butane - In addition to its normal scientific meaning, a mixture mainly of butanes that ordinarily may contain some propane or pentanes plus.

Calibration - Procedures or operations that establish the accuracy of the values indicated by a measuring device as compared to the values indicated by a calibration instrument that has accuracy traceable to a suitable measuring standard. Adjustments are made, if required, to the measuring device to make it comparable to the calibration instrument. May also refer to the procedure used to determine the volume of a prover.

Central Processing Facility - A battery system or arrangement of tanks or other surface equipment receiving the effluent of one or more wells or a satellite prior to delivery to market or other disposition; may include equipment or devices for separating the effluents into crude bitumen, gas, or water for the injection and distribution of air, steam, gas, hydrocarbon, or other materials, for water treatment and recycling, and for measurement but does not include a processing plant.

Clean Oil - Oil with 0.5% S&W or less.

Common Crown or Freehold Royalty - When all the wells in a battery are produced under Crown mineral leases and the Crown receives the same royalty rate for each well, or when

under leases granted by one Freehold mineral holder, the Freehold mineral holder receives the same royalty rate for each well. If there is more than one Freehold mineral holder for the wells in a battery, the total royalty rate for each well is the same.

Common Ownership - All wells in a battery belong to the same working interest participant, or if there is more than one working interest participant, each working interest participant has the same percentage interest in each well in the battery.

Compressor Station - An installation of service equipment that receives natural gas from a well, battery, or gathering system prior to delivery to market or other disposition and is intended to maintain or increase the flowing pressure of the gas; includes any equipment for measurement.

Condensate - A mixture mainly of pentanes and heavier hydrocarbons that may be contaminated with sulphur compounds, that is recovered or is recoverable at a well from an underground reservoir and that may be gaseous in its virgin reservoir state but is liquid at the conditions under which its volume is measured or estimated.

Confidence Level - The degree of confidence that may be placed on an estimated range of uncertainty.

Crude Bitumen - A naturally occurring viscous mixture, mainly of hydrocarbons heavier than pentane, that may contain sulphur compounds and that in its naturally occurring viscous state will not flow to a well.

Crude Oil - A mixture mainly of pentanes and heavier hydrocarbons that may be contaminated with sulphur compounds, that is recovered or is recoverable at a well from an underground reservoir, and that is liquid at the conditions under which its volume is measured or estimated; includes all other hydrocarbon mixtures so recovered or recoverable except raw gas, condensate, or crude bitumen.

Custom Treating Plant - A system or arrangement of tanks and other surface equipment receiving oil/water emulsion exclusively by truck for separation prior to delivery to market or other disposition.

Dehydrator - An apparatus or a process designed and used to remove water from raw gas.

Delivery Point - The point at which the delivery of oil or gas production from a battery or facility is measured. The volumes determined at this point are typically used in royalty calculations, such as sales, cross border, gas plant to facility, or gas plant to gas plant meters.

Delivery Point Measurement - The type of measurement required at a delivery point.

Electronic Flow Measurement (EFM) - Any flow measurement and related system that collects data and performs flow calculations electronically.

Emulsion - A combination of two immiscible liquids, or liquids that do not mix together under normal conditions.

Enhanced Recovery - The increased recovery from a pool achieved by artificial means or by the application of energy extrinsic to the pool; such artificial means or application includes pressuring, cycling, pressure maintenance, or injection to the pool of a substance or form of

energy but does not include the injection in a well of a substance or form of energy for the sole purpose of
- aiding in the lifting of fluids in the well, or
- stimulating the reservoir at or near the well by mechanical, chemical, thermal, or explosive means.

Equilibrium vapour pressure (EVP) - The pressure at which a liquid and its vapour are in equilibrium at a given temperature. When a hydrocarbon liquid has an EVP above the standard pressure (101.325 kPa at 15°C), the EVP at 15°C is the pressure base.

Error - The difference between true and observed values.

Error (random) - An error that varies in an unpredictable manner when a large number of measurements of the same variable are made under effectively identical conditions.

Error (spurious) - A gross error in procedure (for example, human errors or machine malfunctions).

Error (systematic) - An error that in the course of a number of measurements made under the same conditions on material having the same true value of a variable either remains constant in absolute value and sign or varies in a predictable manner. Systematic errors result in a bias.

Ethane - In addition to its normal scientific meaning, a mixture mainly of ethane that ordinarily may contain some methane or propane.

Exception - Circumstances under which if specific criteria are met or approval is granted, measurement devices or procedures are allowed to deviate within specified limits from base measurement requirements.

Facility - Any building, structure, installation, equipment, or appurtenance over which the ERCB has jurisdiction and that is connected to or associated with the recovery, development, production, handling, processing, treatment, or disposal of hydrocarbon-based resources or any associated substances or wastes; includes, without limitation, a battery, processing plant, gas plant, oilfield waste management facility, central processing facility as defined in the *Oil Sands Conservation Regulation* (AR 76/88), compressor, dehydrator, separator, treater, custom treating plant, produced water-injection plant, produced water disposal plant, miscible flood injection plant, satellite, or any combination of any of these, but does not include a well, pipeline as defined in the *Pipeline Act*, mine site or processing plant as defined in the *Oil Sands Conservation Regulation* (AR 76/88), or mine site or coal processing plant as defined in the *Coal Conservation Act*.

Gas - Raw gas or marketable gas or any constituent of raw gas, condensate, crude bitumen, or crude oil that is recovered in processing and that is gaseous at the conditions under which its volume is measured or estimated.

Gas Battery - A system or arrangement of surface equipment receiving the effluent from one or more gas wells that provides separation, measurement, dehydration, dew point control, compression, or other gas handling functions prior to the delivery to market or other disposition; does not include gas processing equipment.

Gas equivalent factor (GEF) - A factor based on the composition of a hydrocarbon liquid mixture that is used to convert the same hydrocarbon liquid mixture to its equivalent gas volume. This factor is mixture dependent and not a constant for all mixtures.

Gas equivalent volume (GEV) - The volume of gas ($10^3 m^3$) that would result from converting 1 m^3 of liquid into a gas by applying a GEF to the liquid volume.

Gas Fractionation Plant - An arrangement of equipment to reprocess an NGL inlet into one or more in-stream components.

Gas Gathering System - A reporting entity that may consist of pipelines used to move gas production from oil batteries, gas batteries, and/or other facilities to another facility (usually a gas plant); may include compressors, line heaters, dehydrators, measurement, and other equipment.

Gas-in-Solution (GIS) - Gas dissolved in an oil volume under pressure.

Gas Plant - A system or arrangement of equipment used for the extraction of hydrogen sulphide, helium, ethane, natural gas liquids, or other substances from raw gas. Does not include a wellhead separator, treater, dehydrator, or production facility that recovers less than 2 m^3/day of hydrocarbon liquids without using a liquid extraction process (e.g., refrigeration, desiccant). In addition, does not include an arrangement of equipment that removes small amounts of sulphur (less than 0.1 tonne/day) through the use of non-regenerative scavenging chemicals that generate no hydrogen sulphide or sulphur dioxide.

Gas Well - A well that produces primarily gas from a pool or portion of a pool wherein the hydrocarbon system is gaseous or exhibits a dew point on reduction of pressure, or any well so designated by the ERCB.

Good Production Practice (GPP) - Production of crude oil or raw gas at a rate
- not governed by a base allowable, but
- limited to what can be produced without adversely and significantly affecting conservation, the prevention of waste, or the opportunity of each owner in the pool to obtain his share of production.

Heavy Oil - Crude oil having a density of 920 kg/m^3 or greater at 15°C.

High vapour pressure liquids - Fluids extracted from raw gas that has been processed at a gas facility, such as ethane, propane, butane, NGL, and in some cases pentanes plus.

Hydrocarbon Liquid - A fluid in the liquid state that may consist of one or more of the following: oil, bitumen, condensate, ethane, propane, butane, pentane plus, or other heavier hydrocarbon compounds.

Initial Qualifying Criteria - Criteria that must be met to qualify for an exception. If the initial qualifying criteria have been met and the exception is implemented, it may remain in place indefinitely, as long as the wells do not meet any of the revocation clauses and there are no physical additions to the battery or facility (e.g., new wells or zones). If additions or changes are made to the battery or facility, the initial qualifying criteria must be met for all the wells or zones added to the battery or facility for the exception to remain in place.

Injection/Disposal Facility - A system or arrangement of surface equipment associated with the injection or disposal of any substance through one or more wells.

Innage Gauge - The depth of liquid in a tank as measured from the surface of the liquid to the tank bottom or to a fixed datum plate.

In Situ Operation
- a scheme or operation ordinarily involving the use of well production operations for the recovery of crude bitumen from oil sands, or
- a scheme or operation designated by the ERCB as an in situ operation, but does not include a mining operation.

Lease Automatic Custody Transfer (LACT) - defined by API MPMS, Chapter 1.

Licensee - The holder of a licence according to the records of the ERCB; includes a trustee or receiver-manager of property of a licensee.

Liquefied petroleum gas (LPG) - LPG consists primarily of propane (C3) and butane (C4) in a mixture or essentially pure form, with minor components ranging from ethane (C2) to normal hexane (C6). It is produced either as a by-product of natural gas processing or during refining and processing operations.

Liquid/Gas Ratio (LGR) - A ratio calculated by dividing the total water and/or condensate test volumes by the measured test gas volume.

Load Fluids - Any hydrocarbon-and/or water-based fluids used at any stage in the life of a well (completion, servicing, regular operation, or abandonment). It includes fluids injected into a flow line between a well and the battery to which it produces (e.g., hot oil, dewaxing chemicals).

Load Oil - Hydrocarbon-type fluids used as load fluid, including crude oil, condensate, refined oils, and oil-based or oil-soluble chemicals.

Load Water - Water-type fluids used as load fluid, including produced/fresh/brackish water and water-based or water-soluble chemicals.

Master Meter - A meter of known accuracy that is connected in series with another meter for the purpose of checking the accuracy of that meter and providing a meter factor.

Maximum Uncertainty of Monthly Volume - Relates to the limits applicable to equipment and/or procedures used to determine the total monthly volume.

Mean - A value in the middle of two extremes (the two values farthest apart in a group of values).

Measured Gas Source(s) - Single-phase measured gas source(s) downstream of separation and removal of liquids; also includes the gas equivalent volume (GEV) of measured condensate if the condensate is recombined after measurement with the gas downstream of the separator.

Measured Oil - Oil measured using equipment and/or procedures meeting delivery point measurement requirements and/or uncertainty limits. For emulsion, the delivery point measurement uncertainty limits apply to the total volume determination only.

Measurement - A procedure for determining a value for a physical variable.

Measurement by Difference - Any situation where an unmeasured volume is determined by taking the difference between two or more measured volumes.

Measuring Standard - A device used in calibration or proving that has a known value traceable to national reference standards maintained by the National Research Council in Ottawa, Ontario.

Meter Factor - A dimensionless number used to correct indicated meter volumes to adjusted volumes if the two volumes differ due to operational conditions (e.g., gas entrained in liquids, meter slippage, meter wear). It is not to be confused with the "K Factor," which is used to convert the number of pulses generated by a meter (where electronic pulse technology is the basis of the meter operation) into units of volume. The K Factor is typically determined by the meter manufacturer and does not take into consideration all of the specific operational conditions the meter may be subjected to.

Metering Difference - The volume used to balance, on a monthly basis, any difference that occurs between the measured inlet/receipt volumes and the measured outlet/disposition volumes at a facility.

Methane - In addition to its normal scientific meaning, a mixture mainly of methane that ordinarily may contain some ethane, nitrogen, helium, or carbon dioxide.

Natural Gas Liquid (NGL) - Propane, butanes, or pentanes plus, or a combination of them, obtained from the processing of raw gas or condensate.

Oil - Condensate or crude oil, or a constituent of raw gas, condensate, or crude oil that is recovered in processing, that is liquid at the conditions under which its volume is measured or estimated.

Oil Battery - A system or arrangement of surface equipment receiving the effluent from one or more oil wells that provides separation, treating, measurement, and other oil handling functions prior to the delivery to market or other disposition.

Oilfield Waste – An unwanted substance or mixture of substances that results from the construction, operation, abandonment, or reclamation of a facility, well site, or pipeline, but does not include an unwanted substance or mixture of substances from such a source that is received for storage, treatment, disposal, or recycling at a facility that is regulated by the Department of Environment.

Oil Well - A well that produces primarily liquid hydrocarbons from a pool or portion of a pool wherein the hydrocarbon system is liquid or exhibits a bubble point on reduction of pressure, or any well so designated by the ERCB.

Operator - With respect to a well or facility, a person who
• has control of or undertakes the day-to-day operations and activities at a well or facility, or

- keeps records and submits production reports for a well or facility to the ERCB, whether or not that person is also the licensee or approval holder of the well or facility.

Outage Gauge - The distance from a reference point at the top of a tank to the surface of the liquid. This "gauge" is then subtracted from the full height gauge (from the same reference point) of the tank to determine the depth of the liquid.

Pentanes Plus - A mixture mainly of pentanes and heavier hydrocarbons, which ordinarily may contain some butanes, that is obtained from the processing of raw gas, condensate, or crude oil.

Pipeline - Any pipe or any system or arrangement of pipes wholly within Alberta and whereby oil, gas, or synthetic crude oil or water incidental to the drilling for or production of oil, gas or synthetic crude oil is conveyed, and
- includes all property of any kind used for the purpose of, in connection with, or incidental to the operation of a pipeline in the gathering, transporting, handling, and delivery of oil, gas, synthetic crude oil, or water, but
- does not include any pipe or any system or arrangement of pipes that constitutes a distribution system for the distribution within a community of gas to ultimate consumers.

Pool - A natural underground reservoir containing or appearing to contain an accumulation of oil or gas or both separated or appearing to be separated from any other such accumulation.

PRA - Petroleum Registry of Alberta.

Propane - In addition to its normal scientific meaning, a mixture mainly of propane that ordinarily may contain some ethane or butanes.
Proration - An accounting procedure or system in which the total actual monthly battery production is equitably distributed among the wells in the battery.

Proration Battery - A battery for which all well production is commingled prior to the total battery production volumes being separated and measured (or estimated where appropriate) as single phases. Individual monthly well production volumes are estimated based on periodic well tests and are corrected to the actual monthly volumes through the use of proration factors.

Prover - A device used to collect and determine the volume of a sample of fluid that has passed through a meter. Provers typically use volumetric or gravimetric means to determine the quantity of the sample.

Prover Run - The operation of a prover or master meter whereby a representative volume is sampled and measured, and that sample volume is compared to the volume indicated by a meter through which the same sample has passed to determine a meter factor.

Proving - The procedures or operations whereby a prover volume is compared to an indicated meter volume (both corrected to applicable pressure and temperature conditions). The prover volume divided by the indicated meter volume yields a meter factor. The meter factor is subsequently applied to indicated meter volumes to determine the adjusted or corrected volume. If the meter is connected to an electronic readout, the meter factor may be incorporated into the software such that the indicated meter volume is already corrected to read the adjusted volume. (Care should be taken in such cases not to apply the meter factor again – in such cases the meter factor should be indicated on the tag or label as being 1.0000.)

Raw Gas - A mixture containing methane, other paraffinic hydrocarbons, nitrogen, carbon dioxide, hydrogen sulphide, helium, and minor impurities, or some of these, that is recovered or is recoverable at a well from an underground reservoir and is gaseous at the conditions under which its volume is measured or estimated.

Relative Density of Gas - The ratio of the mass of the gas to the mass of an equal volume of air. It is also referred to as gas gravity or specific gravity of gas.

S&W - "Sediments and water," commonly refers to settled solid and/or semi-solid components and water in tanks and other containment vessels.

Sales Gas - A mixture mainly of methane originating from raw gas, if necessary through the processing of the raw gas for the removal or partial removal of some constituents, and that meets specifications for use as a domestic, commercial, or industrial fuel or as an industrial raw material.

Satellite or Satellite Battery - Surface equipment located between a number of wells and the main battery that is intended to separate and measure the production from each well, after which the fluids are recombined and piped to the main battery for separation/treating, measurement, and storage or delivery.

Segregate - To confine each fluid in a well to its proper pool or flow channel so that it is separate from the fluid in or passing from or to any other pool.

Separator - An unfired apparatus specifically designed and used for separating fluids produced from a well into two or more streams; does not include a dehydrator.
Single Point Measurement Uncertainty - Relates to the limits applicable to equipment and/or procedures used to determine a specific volume at a single measurement point.

Solid - A substance that does not contain free liquids and is not gaseous at standard conditions.

Solution Gas - Volatile hydrocarbons that are dissolved in solution with produced oil or bitumen.

Stock Tank Vapours - The solution gas present in the oil/bitumen storage tanks that may be released from the tanks.

Synthetic Crude Oil - A mixture mainly of pentanes and heavier hydrocarbons, which may also contain sulphur compounds, that is derived from crude bitumen and is liquid at the conditions under which its volume is measured or estimated; includes all other hydrocarbon mixtures so derived.

Tank - A device designed to contain materials produced, generated, and used by the upstream petroleum industry that is constructed of impervious materials to provide structural support; may include such materials as concrete, plastic, fiberglass reinforced plastic, or steel.

Thief - An industry term for a bottom closure, core-type liquid sampling device used to secure samples from storage tanks.

Treater - A fired apparatus specifically designed and used for separating gas and water from crude oil.

Truck Terminal - A system or arrangement of tanks and other surface equipment that receives hydrocarbon liquids by truck for the purpose of delivering those liquids into a pipeline.

Well - An orifice in the ground completed or being drilled
- for the production of oil or gas,
- for injection to an underground formation,
- as an evaluation well or test hole, or
- to or at a depth of more than 150 m for any purpose;

does not include such an orifice for the purpose of discovering or evaluating a solid inorganic mineral that does not or will not penetrate a stratum capable of containing a pool or oil sands deposit.

Well Event - A unique identifier code for an interval, zone, pool, or horizon in a well. A well may be completed in multiple intervals, zones, pools, or horizons, each of which is identified as a well event.

Appendix 4   Water-Cut (S&W) Procedures

Water-cut procedures are divided into three categories and described on the following pages. Different procedures are specified for the three categories to improve accuracy and consistency of the S&W determinations. The use of "mason jars" with measuring tape attached is not acceptable for determining S&W. S&W percentage must be recorded to a minimum of one decimal place.

More detail on S&W determination is in API MPMS, Chapter 10.4: Determination of Water and/or Sediment in Crude Oil by the Centrifuge Method (Field Procedure). The ERCB will consider any procedure that meets API MPMS, 10.4 standards to be in compliance with this directive. It is the responsibility of the licensee/operator to show that its procedure meets the above API standard.

Category 1—for 0 to 10% S&W

Obtain a representative sample of liquid.

Shake the sample container vigorously to mix it before pouring into the centrifuge tubes.

1) Fill each of two tubes with exactly 100 parts (50 ml) of the sample.

2) Fill each tube with the solvent solution (premixed solvent and demulsifier) to the 200-part mark (100 ml).

3) Stopper each tube tightly and invert 10 times.

4) Loosen the stoppers and immerse the tubes in a preheater. Heat the contents to 60°C ±3°C.

5) Stopper each tube tightly and invert 10 times.

6) Place the tubes in the centrifuge machine in a balanced condition and spin for 5 minutes.

7) Immediately after the centrifuge comes to rest, use a thermometer to verify that the sample temperature is within 9°C of the test temperature.

    If sample temperature is within 9°C, go to step 8. If sample temperature is not within 9°C, go back to step 4, raise the temperature, and repeat steps 5, 6, and 7.

8) Read and record the volume of water and sediment at the bottom of each tube.

9) Reheat the tubes to the initial spin temperature and return them, without agitation, to the centrifuge machine. Spin for an additional 5 minutes. Repeat the procedure until two consecutive, consistent readings are obtained.

    For the test to be considered valid, a clear interface must be observed between the oil layer and the separated water. No emulsion should be present immediately above the oil/water interface. A test comprises TWO TUBES of the SAME SAMPLE. Compare the readings of the two tubes. If the difference is greater than one subdivision on the centrifuge tube, the test is invalid and should be repeated.

10) Calculation and reporting:

For 200 ml tubes: the percentage of water and sediment is the average, to three decimal places, of the values read directly from the two tubes.

For 100 ml tubes: read and record the volume of water and sediment in each tube. Add the readings together and report the sum as the percentage of water and sediment.

Example 1 (see Figure A4.1)

100 ml centrifuge tubes	200 ml centrifuge tubes
If reading from each tube is the same:	
Reading from each tube = 0.50 ml	Reading from each tube = 1.00 ml
Water cut = (0.50 + 0.50) ÷ 100 = 1.0 %	Water cut = 1.00 ÷ 100 = 1.0 %
If reading from each tube is not the same:	
Reading from 1st run of each tube = 0.50, 0.60 ml	Reading from 1st run of each tube = 1.00, 1.05 ml
Reading from 2nd run of each tube = 0.50, 0.55 ml	Reading from 2nd run of each tube = 1.00, 1.10 ml
Water cut = (0.50 + 0.60 + 0.50 + 0.55) ÷ 2 ÷ 100 = 1.1%	Water cut = (1.00 + 1.05 + 1.00 + 1.10) ÷ 4 ÷ 100 = 1.0%
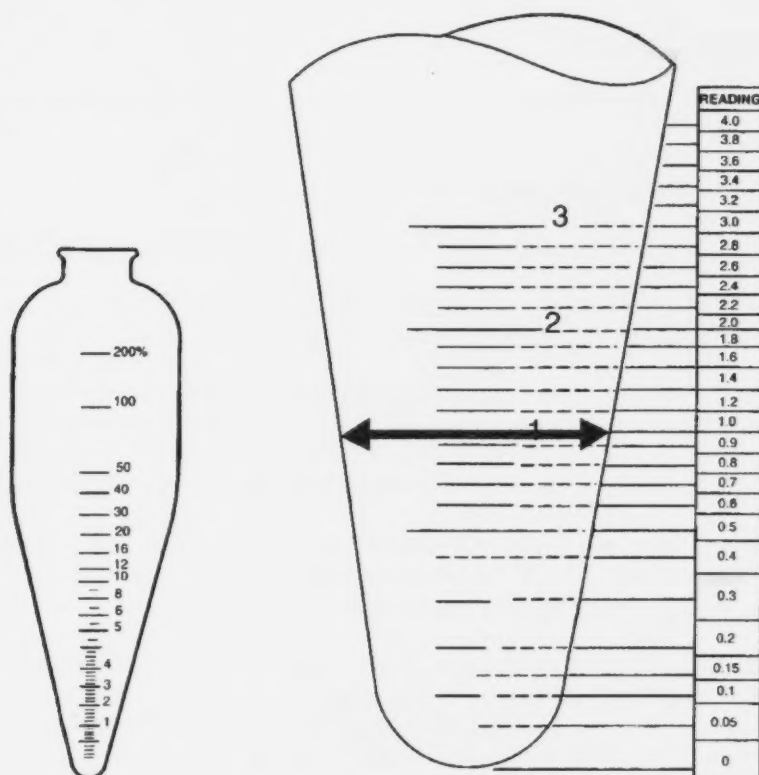
Figure A4.1. Reading a centrifuge tube

Category 2—for 10 to 80% S&W

Obtain the maximum representative sample of liquid feasible (minimum 800 ml).

Transfer the entire sample into an adequately sized graduated cylinder. It may be necessary to wash out the inside of the sample container with a measured volume of solvent to ensure that all of the oil is removed. If this is done, it is necessary to account for the additional amount of solvent added when calculating the water-cut percentage.

Place the graduated cylinder into a heat bath at or above treater temperature (or at or above 60°C if no treater is involved) until the sample temperature and free water fallout have stabilized. A clear oil/water interface must be visible.

Read and record the total volume, the volume of free water, and the volume of oil/emulsion in the graduated cylinder. Calculate the free water percentage as follows:

Percentage of free water = (Volume of free water ÷ Total volume) x 100%

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

Percentage of free water =

Volume of free water ÷ (Total volume – Volume of solvent/demulsifier) x 100%

Draw 100 ml from the oil/emulsion portion in the graduated cylinder and fill each of two 100 ml centrifuge tubes to exactly the 50 ml mark. Add solvent to bring the level in the tubes to exactly the 100 ml mark. The procedures previously outlined for samples with 0 to 10% water cut are to be followed, with the exception that the water-cut readings from both tubes are to be added together, even if they are not the same.

Note that if 200 ml tubes are to be used, a larger initial sample will be required, and if the water-cut readings from both tubes are not the same, the average of both tubes is to be used as the resultant water cut of the oil/emulsion portion.

From the spinning results, calculate the percentage of water remaining in the oil/emulsion portion as follows:

Percentage of water remaining =

Total oil/emulsion volume in cylinder x Water-cut % of oil/emulsion ÷ Total volume

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

Percentage of water remaining =

(Total oil/emulsion volume in cylinder x Water-cut % of oil/emulsion) ÷ (Total volume – Volume of solvent/demulsifier)

Calculate the total water-cut percentage as follows:

Total water-cut % = % free water + % water remaining

Example 2 (see Figure A4.2)

1000 ml graduated cylinder

% of free water = 600 ml ÷ 900 ml x 100% = **66.7%**

% of water remaining = 300 ml x 10%* ÷ 900 ml = **3.3%**

Total Water-cut % = 66.7% + 3.3%= **70.0%**

* Water cut of oil portion determined by spinning samples
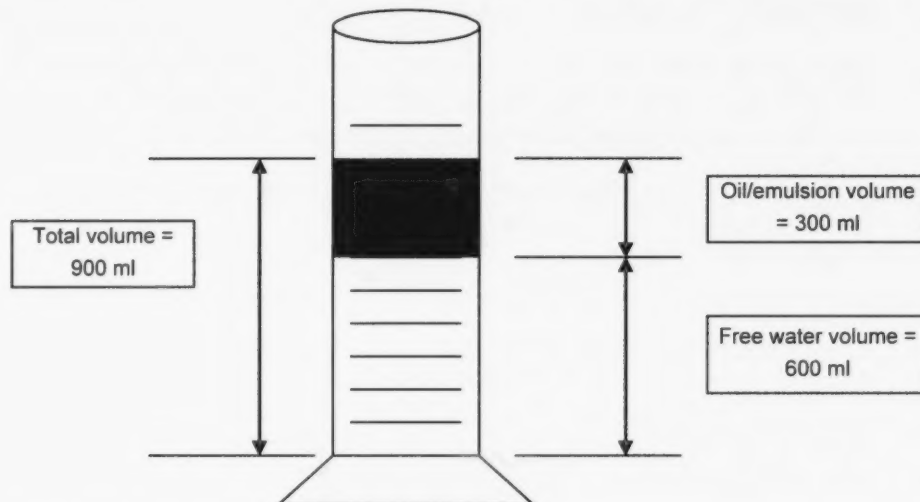
Figure A4.2. Water-cut % from 10 to 80%

Category 3—for 80 to 100% S&W

Obtain the maximum representative sample of liquid feasible (minimum 800 ml).

Transfer the entire sample into an adequately sized graduated cylinder. It may be necessary to wash out the inside of the sample container with a measured volume of solvent to ensure that all of the oil is removed. If this is done, it is necessary to account for the additional amount of solvent added when calculating the water-cut percentage.

Place the graduated cylinder into a heat bath at or above treater temperature (or at or above 60°C if no treater is involved) until the sample temperature and free water fallout have stabilized. A clear oil/water interface must be visible. A narrow-necked graduated cylinder should be used to improve accuracy in sample measurement when the water cut is above 90% (see Figure A4.4).

Read and record the total volume and the volume of free water in the graduated cylinder. If no solvent or demulsifier has been added to the sample, calculate the water-cut percentage as follows:

Water-cut % = Volume of free water ÷ Total volume x 100%

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

Water-cut % = Volume of free water ÷ (Total volume - Volume of solvent/demulsifier) x 100%

The water content of the oil/emulsion portion in the graduated cylinder does not have to be determined, due to the limited amount of the oil/emulsion portion of the sample available at these high water contents. However, if there is enough oil/emulsion volume, the licensee may choose to use the same procedure as that described for the 10 to 80% S&W, with the option to centrifuge only one sample.

Example 3 (see Figure A4.3)

1000 ml graduated cylinder

Water-cut % = 900 ml ÷ 1000 ml x 100%
        = 90.0%

Oil/emulsion volume = 100 ml

Total volume = 1000 ml

Free water volume = 900 ml
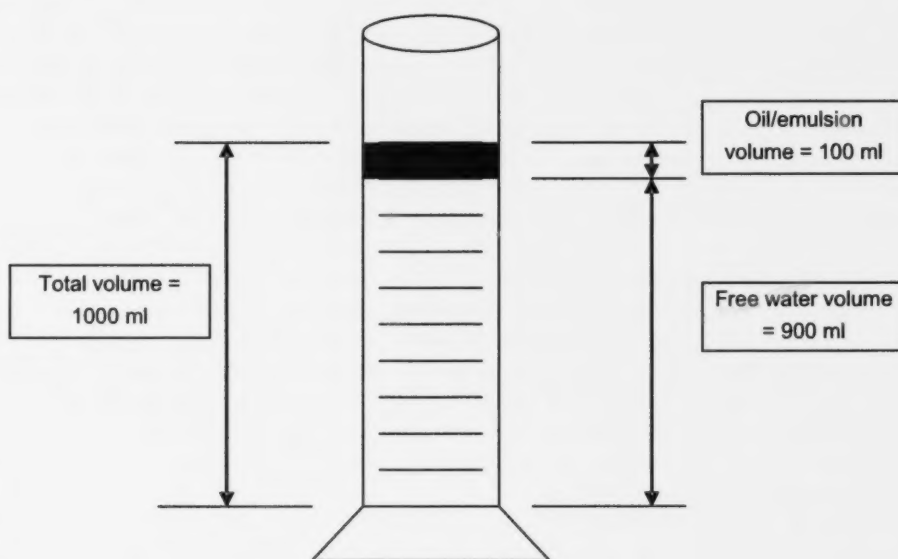
Figure A4.3. Water-cut % over 80%
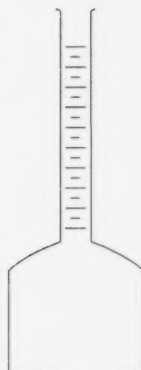
Figure A4.4. Narrow-necked graduated cylinder

Appendix 5 On-site Analytical Techniques for H$_2$S Measurement

On-site measurement of H$_2$S in natural gas streams can be accomplished by several different methods. The appropriate method should be selected with an understanding of the benefits and limitations of each method.

Length of Stain Tubes (GPA Standard 2377-05): For concentrations below 1500 ppm, the most convenient and economical choice is the use of a "length of stain tube." These devices can suffer from some interference, affecting both the precision and the accuracy of the measurements. Nonetheless, for many purposes this technique can provide H$_2$S measurements of a suitable quality. The understanding is that the measurement uncertainty is potentially less than the risk of H$_2$S degradation if a laboratory method were employed. If the most accurate measurements are required, a second sample can be collected in a suitably inert container and returned to a laboratory for prompt analysis.

Tutweiler Titration (GPA Standard C-1): This technique is the method of choice for on-site analysis when the concentration of H$_2$S is greater than 1500 ppm. The Tutweiler titration can provide accurate measurements of H$_2$S using suitably calibrated glassware and chemicals. Operator skill and proper recording of temperatures and barometric pressure are also key elements for this technique.

Instrumental (in-lab) Analytical Techniques for H$_2$S Measurement

Gas Chromatography with Sulphur Selective Detection (ASTM D-5504-01): Sulphur selective detectors can be coupled with gas chromatographs to achieve a low detection limit for H$_2$S and other sulphur compounds, such as mercaptans, sulphides, and disulphides. These instruments are ideal for low concentrations ranging from sub ppm up to several thousand ppm. The sulphur selective detectors are much less susceptible to hydrocarbon interferences and can also identify other sulphur-containing compounds in addition to H$_2$S. Suitable sulphur selective detectors are sulphur chemiluminescence detectors (SCD) and pulsed flame photometric detectors (PFPD).

Gas Chromatography with Thermal Conductivity Detection: Thermal conductivity detectors can be coupled with gas chromatographs to analyze for intermediate to high levels of H$_2$S. H$_2$S can be adequately resolved from hydrocarbon components to allow for specific detection. The columns selected for this type of analysis must offer a good balance between high resolution (specificity of H$_2$S) and low adsorption of H$_2$S. Detection limits for H$_2$S levels as low as 300 ppm can be achieved under the right conditions, and the method can also be calibrated for values approaching 100 per cent H$_2$S. The analytical range for these systems should not exceed the linear range of the column and detector combination. Therefore, acceptable calibration ranges must yield a linear calibration curve (minimum 4 points) with an R-squared value of no less than 0.99.

Appendix 6   Gas Equivalent Volume Determination

Liquid Analysis Example

Component	Volume Fractions	Mole Fractions	Mass Fractions
N2	0.0006	0.0019	0.0008
CO2	0.0081	0.0158	0.0109
H2S	0	0	0
C1	0.0828	0.1617	0.0405
C2	0.1117	0.1462	0.0687
C3	0.1275	0.1533	0.1056
IC4	0.0394	0.0398	0.0362
NC4	0.0891	0.0935	0.0849
IC5	0.0483	0.0436	0.0492
NC5	0.0540	0.0493	0.0556
C6	0.0765	0.0614	0.0835
C7	0.0880	0.0678	0.1054
C8	0.0827	0.0589	0.1032
C9	0.0570	0.0368	0.0726
C10	0.0363	0.0222	0.0480
C11	0.0225	0.0131	0.0305
C12+	0.0755	0.0347	0.1044
TOTAL	1.0000	1.0000	1.0000

Properties of C5+ & C7+ portion of sample					
	Mol. Fractions	Wt. Fractions	Liq. Vol. Fractions	Mol. Wt. (kg/kmol)	Absolute Density (AD) (kg/m^3)
C5+	0.3878	0.6524	0.5408	107.7	739.33
C7+	0.2335	0.4641	0.3620	127.2	785.29

1 Gas Equivalent Factor by Volume Fraction Calculation

GEF = Total (Pseudo 10^3 m^3 Gas / m^3 Liquid)

Condensate Stream : _____

Component	Vol. Fraction Liquid Analysis		10^3 m^3 Gas / m^3 Liquid @ 101.325 kPa & 15°C		Pseudo 10^3 m^3 Gas / m^3 Liquid
N2		x	0.68040	=	
CO2		x	0.44120	=	
H2S		x	0.55460	=	
C1		x	0.44217	=	
C2		x	0.28151	=	
C3		x	0.27222	=	
IC4		x	0.22906	=	
NC4		x	0.23763	=	
IC5		x	0.20468	=	
NC5		x	0.20681	=	
C6		x	0.18216	=	
C7		x	0.16234	=	
C8		x	0.14629	=	
C9		x	0.13303	=	
C10		x	0.12194	=	
				Total =	

Note: For C5+, C6+, or C7+ Sample:

Properties of C5+, C6+, or C7+ sample @ 15°C

AD = [   ]
Mol. Wt. = [   ]

10^3 m^3 Gas / m^3 Liquid = 23.645 (m^3/kmol) x AD (kg/m^3) / Mol. Wt. (kg/kmol)

= [   ] Input this factor to the table above for C5+, C6+, or C7+

GEF = Total (10^3 m^3 Gas / m^3 Liquid)

= [   ] (10^3 m^3 Gas / m^3 Liquid)

Example 1

Gas Equivalent Factor by Volume Fraction Calculation

GEF = Total (Pseudo 10^3 m^3 Gas / m^3 Liquid)

Condensate Stream : _____

Component	Volume Fraction Liquid Analysis		10^3m^3 Gas / m^3 Liquid @ 101.325 kPa & 15°C		Pseudo 10^3 m^3 Gas / m^3 Liquid
N2	0.0006	x	0.68040	=	0.0004
CO2	0.0081	x	0.44120	=	0.0036
H2S	0	x	0.55460	=	
C1	0.0828	x	0.44217	=	0.0366
C2	0.1117	x	0.28151	=	0.0314
C3	0.1275	x	0.27222	=	0.0347
IC4	0.0394	x	0.22906	=	0.0090
NC4	0.0891	x	0.23763	=	0.0212
IC5	0.0483	x	0.20468	=	0.0099
NC5	0.0540	x	0.20681	=	0.0112
C6	0.0765	x	0.18216	=	0.0139
C7+	0.3620	x	0.14598	=	0.0528
		x		=	
	1.0000			Total =	0.2247

Note: For C7+ Sample:

Properties of C7+ sample @ 15 °C

AD = 785.29
Mol. Wt. = 127.2

10^3 m^3 Gas / m^3 Liquid = 23.645 (m^3/kmol) x AD (kg/m^3) / Mol. Wt. (kg/kmol) / 1000 (m^3/10^3 m^3)

= 0.14598 Input this factor to the table above for C7+

GEF = Total (Pseudo 10^3 m^3 Gas / m^3 Liquid)

= 0.2247 (10^3 m^3 Gas / m^3 Liquid)

2 Gas Equivalent Factor by Mole Fraction Calculation

GEF = 23.645 (m³/kmol) / Total (Pseudo m³/kmol) / 1000 (m³/10³ m³)

Condensate Stream : _____

Component	Mol. Fraction Liquid Analysis		10³ m³/kmol @ 101.325 kPa & 15°C		Pseudo 10³ m³/kmol
N2		x	0.03475	=	
CO2		x	0.05359	=	
H2S		x	0.04263	=	
C1		x	0.05348	=	
C2		x	0.08399	=	
C3		x	0.08686	=	
IC4		x	0.10322	=	
NC4		x	0.09950	=	
IC5		x	0.11552	=	
NC5		x	0.11433	=	
C6		x	0.12980	=	
C7		x	0.14565	=	
C8		x	0.16163	=	
C9		x	0.17774	=	
C10		x	0.19391	=	
				Total =	

Note:

Properties of C5+, C6+, or C7+ Sample @ 15°C:

$$AD = \boxed{}$$
$$Mol.\ Wt. = \boxed{}$$

For C5+, C6+, or C7+ Sample:

10³ m³/kmol = Mol. Wt. / AD

$$= \boxed{}$$

GEF = 23.645 (m³/kmol) / Total (Pseudo m³/kmol) / 1000 (m³/10³ m³)

$$= \boxed{}\ (10^3\ m^3\ Gas\ /\ m^3\ Liquid)$$

Example 2

Gas Equivalent Factor by Mole Fraction Calculation

GEF = 23.645 (m^3/kmol) / Total (Pseudo m^3/kmol) / 1000 (m^3/10^3 m^3)

Condensate Stream : _____

Component	Mol. Fraction Liquid Analysis		m^3/kmol @ 101.325 kPa & 15°C		Pseudo m^3/kmol
N2	0.0019	x	0.03475	=	0.0001
CO2	0.0158	x	0.05359	=	0.0008
H2S	0	x	0.04263	=	0
C1	0.1617	x	0.05348	=	0.0086
C2	0.1462	x	0.08399	=	0.0123
C3	0.1533	x	0.08686	=	0.0133
IC4	0.0398	x	0.10322	=	0.0041
NC4	0.0935	x	0.09950	=	0.0093
IC5	0.0436	x	0.11552	=	0.0050
NC5	0.0493	x	0.11433	=	0.0056
C6	0.0614	x	0.12980	=	0.0080
C7+	0.2335	x	0.16198	=	0.0378
		x		=	
		x		=	
	1.0000			Total =	0.1049

Note:
Properties of C7+ sample @ 15 °C:

AD =	785.29
Mol. Wt. =	127.2

For C7+ Fraction:
m^3/kmol = Mol. Wt. / AD

= [0.16198] Input this factor to the table above for C7+

GEF = 23.645 (m^3/kmol) / Total (Pseudo m^3/kmol) / 1000 (m^3/10^3 m^3)

= [0.2254] (10^3 m^3 Gas / m^3 Liquid)

3 Gas Equivalent Factor by Mass Fraction Calculation

Step 1. Calculate Pseudo Volume (L) = Mass Fraction / Liquid Density x 1000 L/m^3

Step 2. Calculate Volume Fraction = Component Pseudo Volume / Total Pseudo Volume

Step 3. Calculate Component Pseudo GEF = Volume Fraction x (10^3 m^3 Gas / m^3 Liquid)

Condensate Stream : _____

Component	Mass Fraction Liquid Analysis		Liquid Density (kg/m^3)		Step 1 Pseudo Volume (L)	Step 2 Volume Fraction		10^3 m^3 Gas / m^3 Liquid @ 101.325 kPa & 15°C		Step 3 Pseudo 10^3 m^3 Gas / m^3 Liquid
N2		/	806.10	=			x	0.68040	=	
CO2		/	821.22	=			x	0.44120	=	
H2S		/	799.40	=			x	0.55460	=	
C1		/	300.00	=			x	0.44217	=	
C2		/	358.00	=			x	0.28151	=	
C3		/	507.67	=			x	0.27222	=	
IC4		/	563.07	=			x	0.22906	=	
NC4		/	584.14	=			x	0.23763	=	
IC5		/	624.54	=			x	0.20468	=	
NC5		/	631.05	=			x	0.20681	=	
C6		/	663.89	=			x	0.18216	=	
C7		/	687.98	=			x	0.16234	=	
C8		/	706.73	=			x	0.14629	=	
C9		/	721.59	=			x	0.13303	=	
C10		/	733.76	=			x	0.12194	=	
									Total =	

Note: For C5+, C6+, or C7+ Sample:

Mol. Wt. = [   ]

AD   = [   ]

10^3 m^3 Gas / m^3 Liquid = 23.645 x AD / Mol. Wt. / 1000 m^3/10^3 m^3

= [   ]

GEF = Total (Pseudo 10^3 m^3 Gas / m^3 Liquid)

= [   ]   (10^3 m^3 Gas / m^3 Liquid)

Example 3

Gas Equivalent Factor by Mass Fraction Calculation

Step 1. Calculate Pseudo Volume (L) = Mass Fraction / Liquid Density x 1000 L/m³
Step 2. Calculate Volume Fraction based on total pseudo volume
Step 3. Calculate Component Pseudo GEF = Volume Fraction x (10³m³ Gas / m³ Liquid)

Condensate Stream : _____

Component	Mass Fraction Liquid Analysis		Liquid Density (kg/m³)		Step 1 Pseudo Volume (L)	Step 2 Volume Fraction		10³ m³ Gas / m³ Liquid @ 101.325 kPa & 15°C		Step 3 Pseudo 10³ m³ Gas / m³ Liquid
N2	0.0008	/	806.10	=	0.0010	0.0006	x	0.68040	=	0.0004
CO2	0.0109	/	821.22	=	0.0133	0.0081	x	0.44120	=	0.0036
H2S	0	/	799.40	=			x	0.55460	=	
C1	0.0405	/	300.00	=	0.1350	0.0822	x	0.44217	=	0.0363
C2	0.0687	/	358.00	=	0.1919	0.1168	x	0.28148	=	0.0329
C3	0.1056	/	507.67	=	0.2080	0.1266	x	0.27221	=	0.0345
IC4	0.0362	/	563.07	=	0.0643	0.0391	x	0.22906	=	0.0090
NC4	0.0849	/	584.14	=	0.1453	0.0885	x	0.23763	=	0.0210
IC5	0.0492	/	624.54	=	0.0788	0.0480	x	0.20468	=	0.0098
NC5	0.0556	/	631.05	=	0.0881	0.0536	x	0.20681	=	0.0111
C6	0.0835	/	663.89	=	0.1258	0.0766	x	0.18216	=	0.0140
C7+	0.4641	/	785.29	=	0.5910	0.3598	x	0.14598	=	0.0525
	1.0000				1.6425	1.0000		Total =		0.2251

Note: For C7+ Sample:

Mol. Wt. = 127.2
AD of C7+ liquid = 785.29 (kg/m³)

10³ m³ Gas / m³ Liquid = 23.645 x AD / Mol. Wt. x 999.10 / 1000 (m³/10³ m³)

= 0.14598 Input this factor to the table above for C7+

GEF = Total (Pseudo 10³ m³ Gas / m³ Liquid)

= 0.2251 (10³ m³ Gas / m³ Liquid)

Appendix 7 Calculated Compositional Analysis Examples

Calculated Well Stream Compositional Analysis Example

Step 1: Collect volumetric and compositional data for both gas and liquid phases.

Gas	
Gas Volume ($10^3\ m^3$)	10000.0
Composition	Mole %
N2	1.00
CO2	2.00
H2S	2.40
C1	80.00
C2	8.00
C3	3.00
IC4	1.00
NC4	1.50
IC5	0.20
NC5	0.50
C6	0.30
C7+	0.10
	100.00

Liquid	
Liquid Volume (m^3)	200.0
Composition	Mole %
N2	0.00
CO2	1.00
H2S	2.00
C1	3.00
C2	4.00
C3	7.00
IC4	10.00
NC4	15.00
IC5	7.00
NC5	11.00
C6	10.00
C7+	30.00
	100.00

Step 2: Convert the condensate liquid volume to a GEV.

A) Convert liquid volume to equivalent gas volume using the condensate gas equivalent factor.

Equation 1: *GEV = Volume of condensate (m^3) x GEF (m^3 gas per m^3 liquid)*

$GEV = 200 \ (m^3) \times 220.12 \ (m^3 \ gas \ per \ m^3 \ liquid) \div 1000 \ (10^3 \ m^3/m^3) = 44.024 \ 10^3 \ m^3$

If the gas equivalent factor is not included with the condensate analysis report, it can be calculated.

Equation 2: *GEF = Absolute Density of Condensate (kg/m^3 @ 15°C) / Molecular weight of the condensate (grams/mole)*

B) Convert the compositional fractions to equivalent gas volumes on a component basis.

Equation 3: *Component gas volume ($10^3 \ m^3$) = [Component mole percent] x [GEV]*

Example: n-pentane equivalent volume:
Volume of condensate = 200 m^3
Gas Equivalent Factor = 220.12
Equivalent n-pentane (NC5) gas volume = [11.0%] x [44.024 $10^3 \ m^3$] = 4843 m^3

Liquid	
Liquid Volume (m^3)	200.0
Composition	Mole %
N2	0.00
CO2	1.00
H2S	2.00
C1	3.00
C2	4.00
C3	7.00
IC4	10.00
NC4	15.00
IC5	7.00
NC5	11.00
C6	10.00
C7+	30.00
	100.00

A →
B →

Liquid	
Gas Equivalent Volume ($10^3 \ m^3$)	44.024
Composition	$10^3 \ m^3$ gas
N2	0.00
CO2	0.44
H2S	0.88
C1	1.32
C2	1.76
C3	3.08
IC4	4.40
NC4	6.60
IC5	3.08
NC5	4.84
C6	4.40
C7+	13.21
	44.02

Step 3:
Add the gas volumes and liquid gas equivalent volumes and normalize
to mole fraction 1 or 100%

Gas			Liquid			Recombined Volume			Recombined Composition	
Gas Volume $(10^3\,m^3)$	10000.0		Gas Equivalent Volume $(10^3\,m^3)$	44.0		Gas Equivalent Volume $(10^3\,m^3)$	10044.0		Gas Equivalent Volume $(10^3\,m^3)$	10044.0
Composition	$10^3\,m^3$ gas		Composition	$10^3\,m^3$ gas		Composition	$10^3\,m^3$ gas		Composition	Mole %
N2	100.0		N2	0.00		N2	100.0		N2	1.00
CO2	200.0		CO2	0.44		CO2	200.4		CO2	2.00
H2S	240.0		H2S	0.88		H2S	240.9		H2S	2.40
C1	8000.0	+	C1	1.32	=	C1	8001.3		C1	79.66
C2	800.0		C2	1.76		C2	801.8		C2	7.98
C3	300.0		C3	3.08		C3	303.1		C3	3.02
IC4	100.0		IC4	4.40		IC4	104.4		IC4	1.04
NC4	150.0		NC4	6.60		NC4	156.6		NC4	1.56
IC5	20.0		IC5	3.08		IC5	23.1		IC5	0.23
NC5	50.0		NC5	4.84		NC5	54.8		NC5	0.55
C6	30.0		C6	4.40		C6	34.4		C6	0.34
C7+	10.0		C7+	13.21		C7+	23.2		C7+	0.23
	10000.0			44.0			10044.0			100.00

Calculated Group Compositional Analysis Example

Step 1:
Collect Volumetric and Compositional data for both gas and liquid phases for all streams. This information is required for all wells.

Step 2:
Mathematically recombine the fluid based on volumetric and compositional data collected in Step 1 for each stream.

Step 3:
Add the recombined fluid volumes on a component basis and normalize to 100%.

Stream 1: Gas			Liquid			Recombined Fluid	
Gas Volume ($10^3 m^3$)	10000		Gas Equivalent Volume ($10^3 m^3$)	800		Gas Equivalent Volume ($10^3 m^3$)	10800
Composition	Mole %		Composition	Mole %		Composition	Mole %
N2	1.14		N2	0.12		N2	1.06
CO2	0.16		CO2	0.08		CO2	0.15
H2S	0.00		H2S	0.00		H2S	0.00
C1	85.31		C1	22.02		C1	80.62
C2	6.44		C2	6.14		C2	6.42
C3	3.77		C3	8.56		C3	4.12
IC4	0.63		IC4	2.62		IC4	0.78
NC4	1.32		NC4	7.11		NC4	1.75
IC5	0.33		IC5	3.66		IC5	0.58
NC5	0.41		NC5	5.73		NC5	0.80
C6	0.26		C6	9.73		C6	0.96
C7+	0.23		C7+	34.23		C7+	2.75
	100.00			100.00			100.00

Stream 2: Gas			Liquid			Recombined Fluid			Total Recombined Fluid	
Gas Volume ($10^3 m^3$)	15000		Gas Equivalent Volume ($10^3 m^3$)	200		Gas Equivalent Volume ($10^3 m^3$)	15200		Gas Equivalent Volume ($10^3 m^3$)	36000
Composition	Mole %		Composition	Mole %		Composition	Mole %		Composition	Mole %
N2	1.00		N2	0.00		N2	0.99		N2	0.76
CO2	2.00		CO2	1.00		CO2	1.99		CO2	1.44
H2S	2.40		H2S	2.00		H2S	2.39		H2S	1.01
C1	80.00		C1	3.00		C1	78.99		C1	82.37
C2	8.00		C2	4.00		C2	7.95		C2	6.95
C3	3.00		C3	7.00		C3	3.05		C3	2.94
IC4	1.00		IC4	10.00		IC4	1.12		IC4	0.79
NC4	1.50		NC4	15.00		NC4	1.68		NC4	1.37
IC5	0.20		IC5	7.00		IC5	0.29		IC5	0.32
NC5	0.50		NC5	11.00		NC5	0.64		NC5	0.54
C6	0.30		C6	10.00		C6	0.43		C6	0.47
C7+	0.10		C7+	30.00		C7+	0.49		C7+	1.04
	100.00			100.00			100.00			100.00

Stream 3: Gas			Liquid			Recombined Fluid	
Gas Volume ($10^3 m^3$)	10000		Gas Equivalent Volume ($10^3 m^3$)	0		Gas Equivalent Volume ($10^3 m^3$)	10000
Composition	Mole %		Composition	Mole %		Composition	Mole %
N2	0.10		N2	0.00		N2	0.10
CO2	2.00		CO2	0.00		CO2	2.00
H2S	0.00		H2S	0.00		H2S	0.00
C1	89.40		C1	0.00		C1	89.40
C2	6.00		C2	0.00		C2	6.00
C3	1.50		C3	0.00		C3	1.50
IC4	0.30		IC4	0.00		IC4	0.30
NC4	0.50		NC4	0.00		NC4	0.50
IC5	0.08		IC5	0.00		IC5	0.08
NC5	0.10		NC5	0.00		NC5	0.10
C6	0.01		C6	0.00		C6	0.01
C7+	0.01		C7+	0.00		C7+	0.01
	100.00			0.00			100.00

Calculated Single Compositional Analysis (from Two Samples)

Step 1: Collect spot samples and record the metered volumes associated with each sample.

Step 2: Calculate individual component volumes by multiplying the individual component mole fractions or percentage values by the associated metered volumes.

Example: Gas Sample #1, Calculation of methane volume
Total Volume = 10000 $10^3 m^3$
Methane = 80.00 mole %
Methane Volume = 10000.0 $10^3 m^3$ X 0.8000 = 8000.0 $10^3 m^3$

Step 3: Normalization: Individual component volumes are summed. The individual component volumes are then be divided into the total to create a normalized (calculated) compositional value.

Example: Ethane (C2), Calculation of Mole %
Gas Sample #1, C2 volume: 800 $10^3 m^3$
Gas Sample #2, C2 volume: 560 $10^3 m^3$
Combined, C2 volume: 1360 $10^3 m^3$
Total gas volume: 18000 $10^3 m^3$
Calculated C2 concentration = 1360.0 $10^3 m^3$ / 18000 $10^3 m^3$ = 7.56 mole %

Gas Sample #1		
Gas Volume ($10^3 m^3$) = 10000.0		
Component	Mole %	$10^3 m^3$ gas
N2	1.00	100.0
CO2	2.00	200.0
H2S	2.40	240.0
C1	80.00	8000.0
C2	8.00	800.0
C3	3.00	300.0
IC4	1.00	100.0
NC4	1.50	150.0
IC5	0.20	20.0
NC5	0.50	50.0
C6	0.30	30.0
C7+	0.10	10.0
	100.00	10000.0

+

Gas Sample #2		
Gas Volume ($10^3 m^3$) = 8000.0		
Component	Mole %	$10^3 m^3$ gas
N2	0.60	48.0
CO2	2.00	160.0
H2S	1.50	120.0
C1	83.00	6640.0
C2	7.00	560.0
C3	2.50	200.0
IC4	1.00	80.0
NC4	1.40	112.0
IC5	0.18	14.4
NC5	0.45	36.0
C6	0.28	22.4
C7+	0.09	7.2
	100.00	8000.0

=

Calculated Single Compositional Analysis		
Gas Volume ($10^3 m^3$) = 18000.0		
Component	Calculated Mole %	$10^3 m^3$ gas
N2	0.82	148.0
CO2	2.00	360.0
H2S	2.00	360.0
C1	81.33	14640.0
C2	7.56	1360.0
C3	2.78	500.0
IC4	1.00	180.0
NC4	1.46	262.0
IC5	0.19	34.4
NC5	0.48	86.0
C6	0.29	52.4
C7+	0.10	17.2
	100.00	18000.0

Appendix 8   Blending Shrinkage Calculation Example

	API MPMS Chapter 12.3, Section 5.3, Eq. 3			
% Shrinkage = 26900 * C * (100 - C) ^ 0.819 * (1/dL - 1/dH) ^ 2.28			dL = Weighted Average Diluent Density	
C = Diluent Volume / (Bitumen Volume + Diluent Volume)			dH = Weighted Average Crude Density	
Inputs	Volume (m³)	Density (kg/m³)		Volume %
Light Component	800.0	700.0	Light Component % in Blend	25.81%
Heavy Component	2300.0	963.0	Heavy Component % in Blend	74.19%
C = 25.81%				
1/dL = 0.00143				
1/dH = 0.00104	% Shrinkage = 0.3993%			

For blending 2,300 m³ of 963 kg/m³ oil and 800 m³ of 700 kg/m³ diluent:

Shrinkage volume = 0.3993% x (800 + 2300) = 12.38 m³